Figure 1

Red blood cell lysis in donor A after incubation with PMAA-Na in RPMI

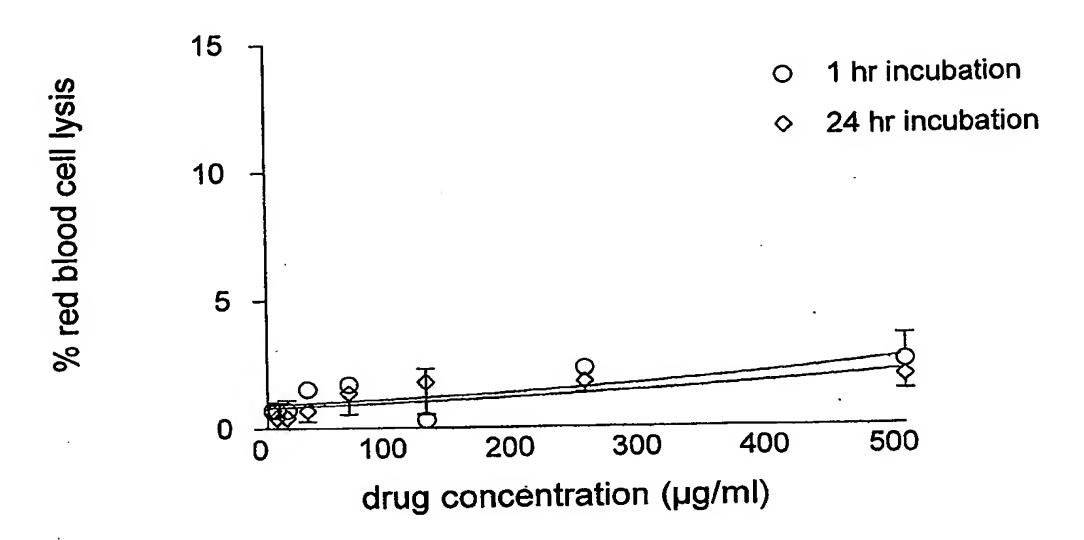


Figure 1a

Red blood cell lysis in donor C after incubation with PMAA-Na in RPMI

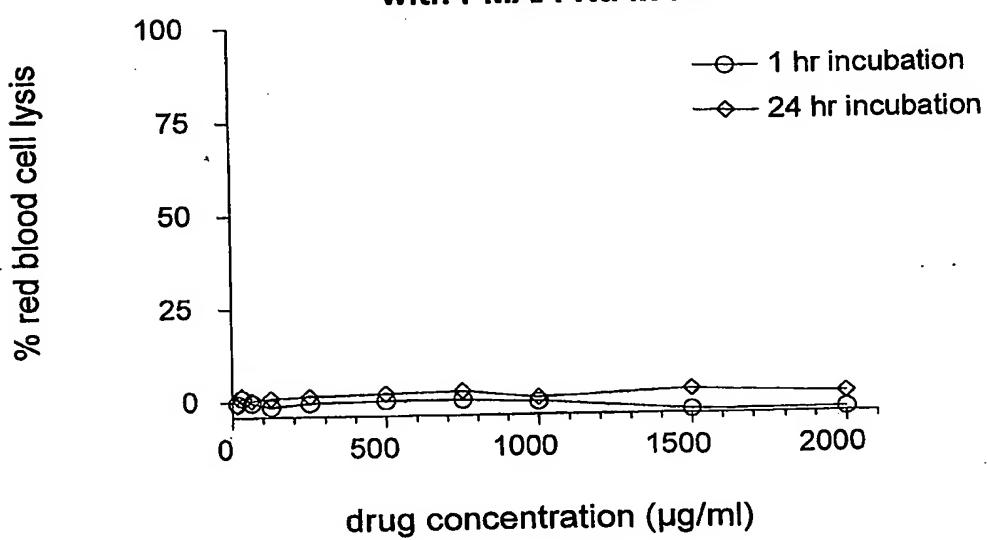


Figure 1b

Figure 1 cont.

Red blood cell lysis in donor B after incubation with PMAA-Na in RPMI

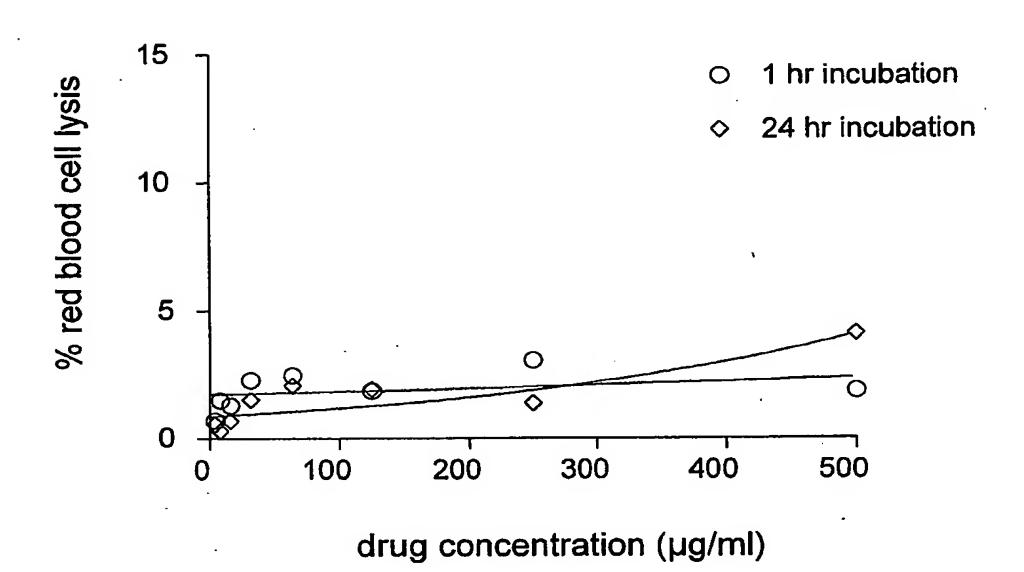
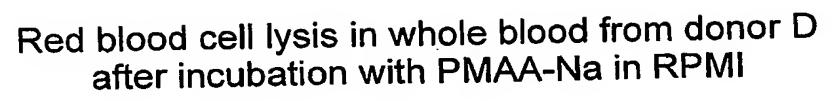


Figure 1c

Figure 2



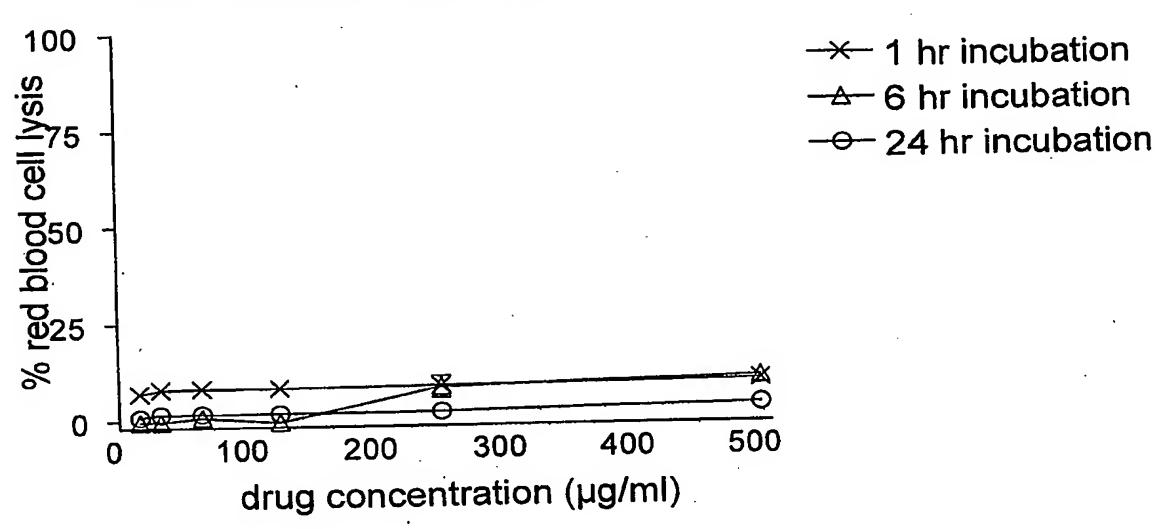


Figure 3

Lack of toxicity of PMAA-Na on primary human monocyte-derivedmacrophages

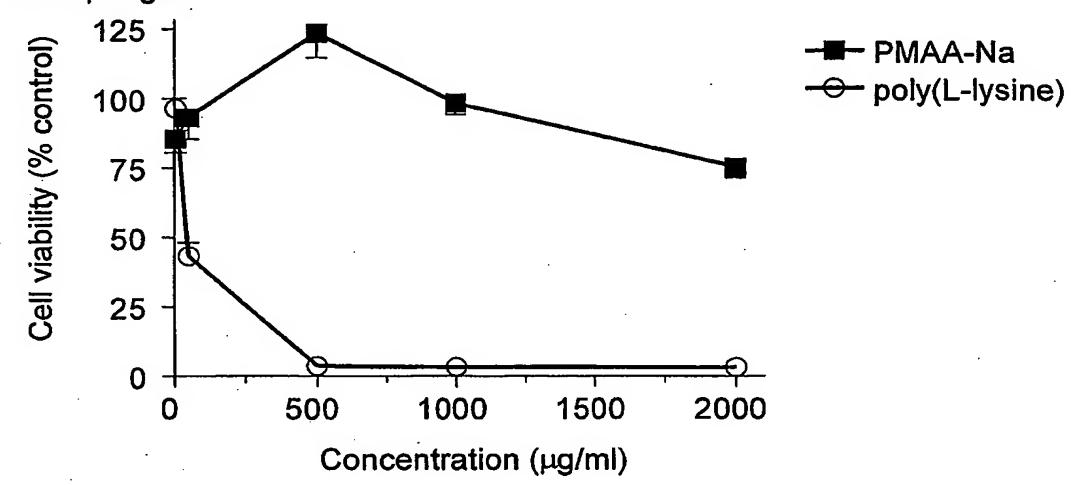


Figure 3a

Lack of toxicity of PMAA-Na on primary human peritoneal macrophages

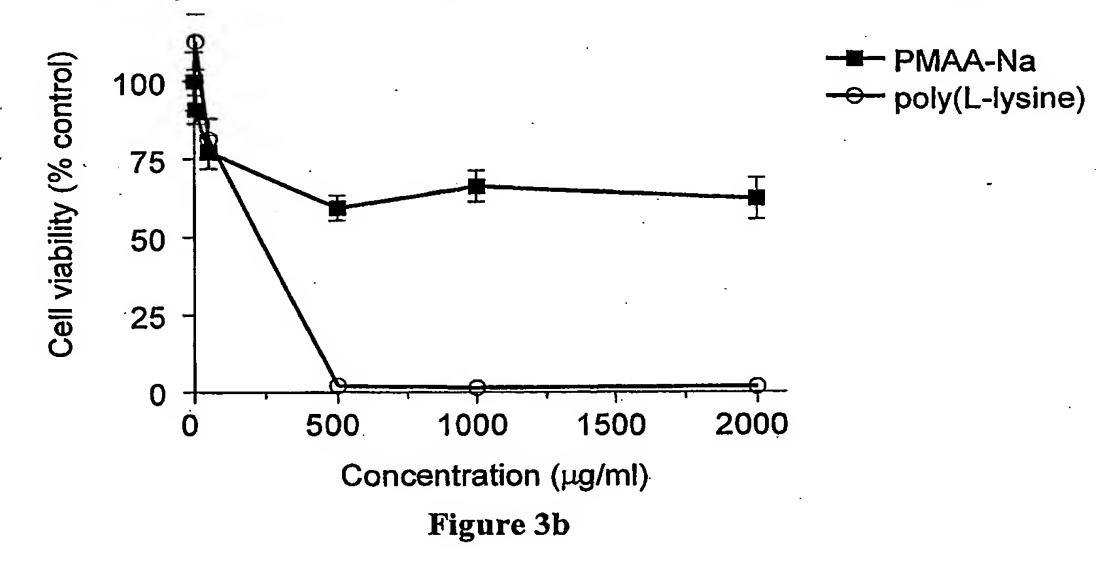


Figure 4

Release of MIP-1 β from human peritoneal macrophages by endotoxin free PMAA-Na (500 μ g/ml) after 36 h from a single donor A

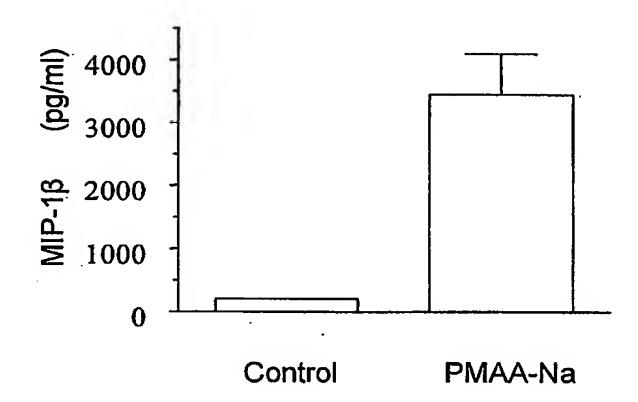


Figure 4a

Release of MIP-1 β from human peritoneal macrophages by endotoxin free PMAA-Na (500 $\mu g/ml$) after 36 h from a single donor B

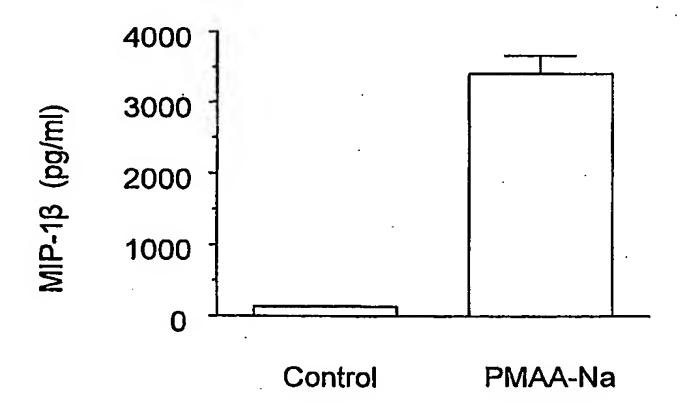


Figure 4b

Figure 4 cont.

Release of MIP-1β from human peritoneal macrophages by endotoxin free PMAA-Na (500 µg/ml) after 36 h from a single donor C

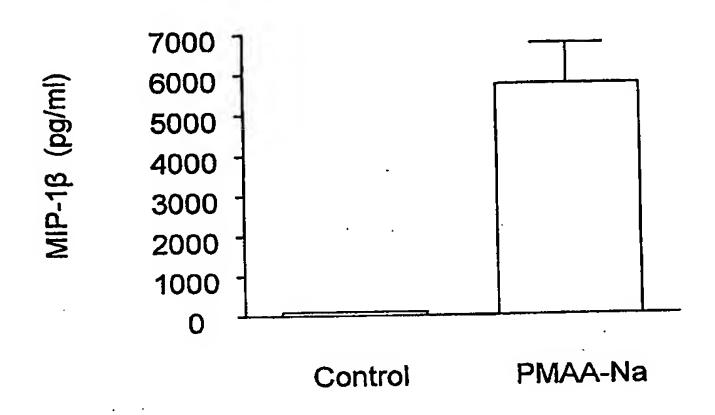


Figure 4c

Figure 5

Release of TNFα from human peritoneal macrophages by endotoxin free PMAA-Na (2,000 μg/ml) after 36 h from a single donor A

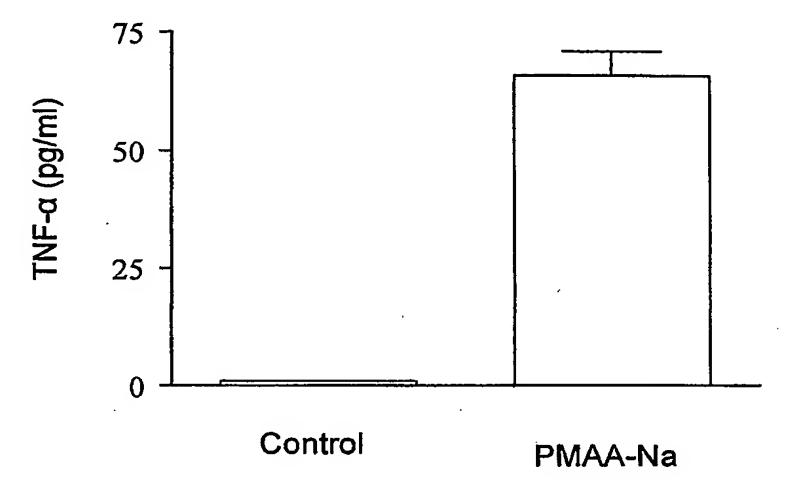


Figure 5a

Release of TNFα from human peritoneal macrophages by endotoxin free PMAA-Na (500 μg/ml) after 36 h from a single donor B

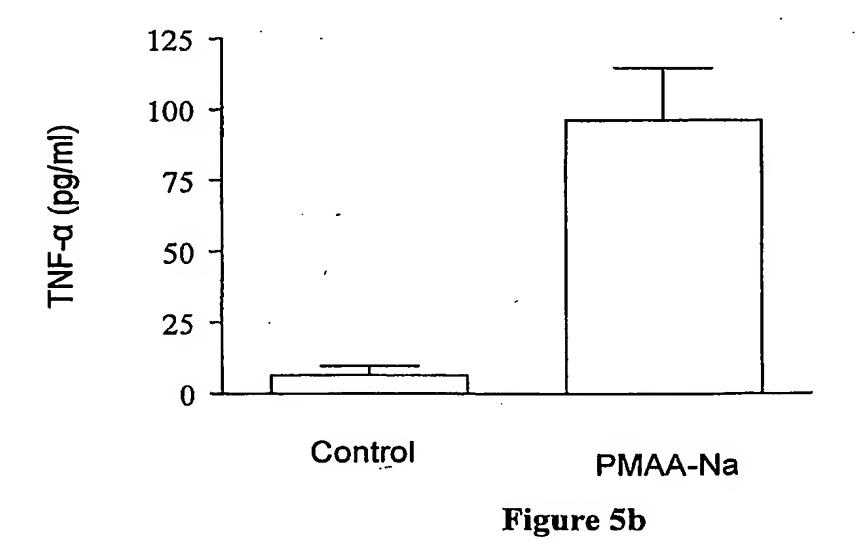
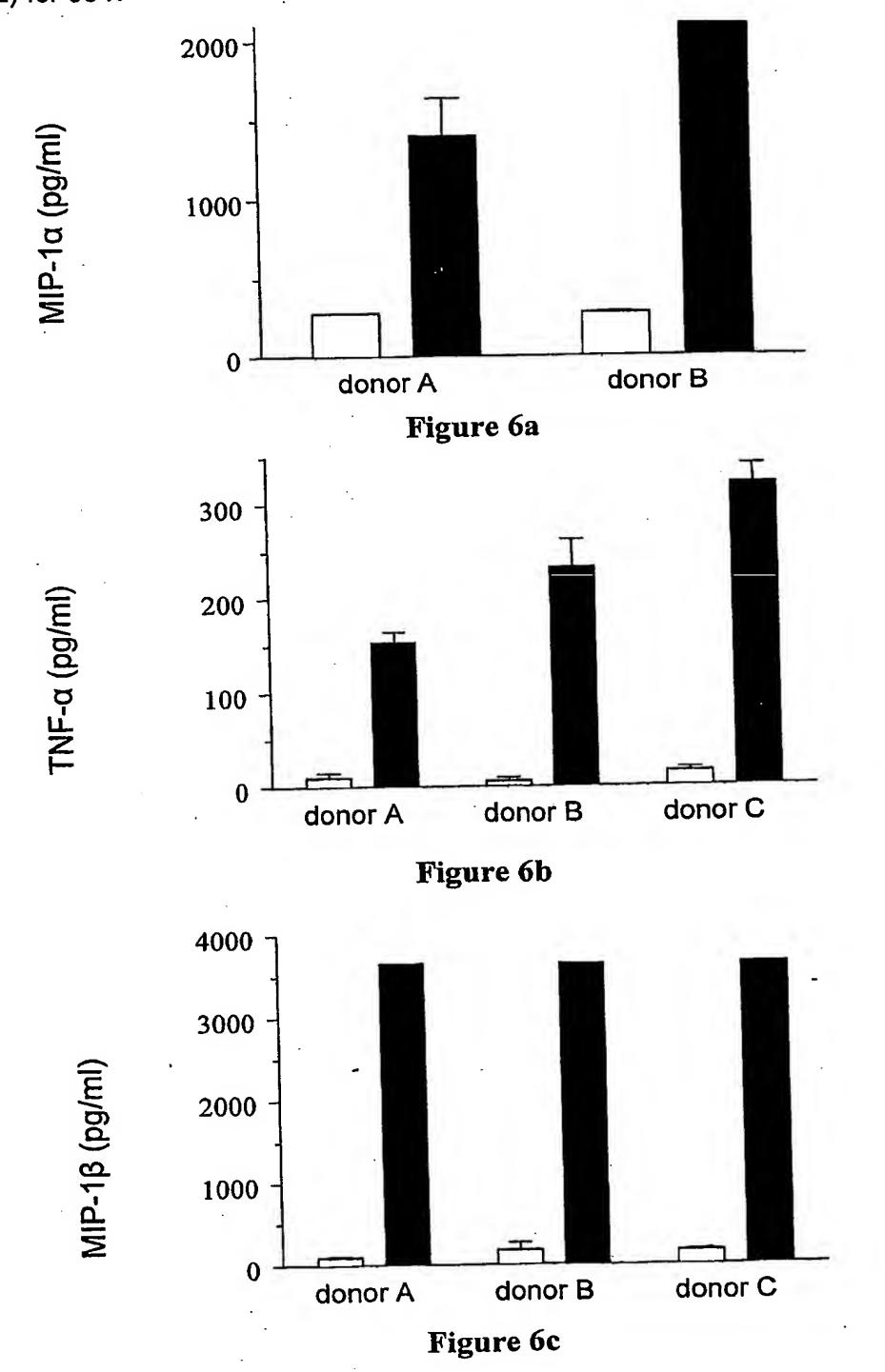
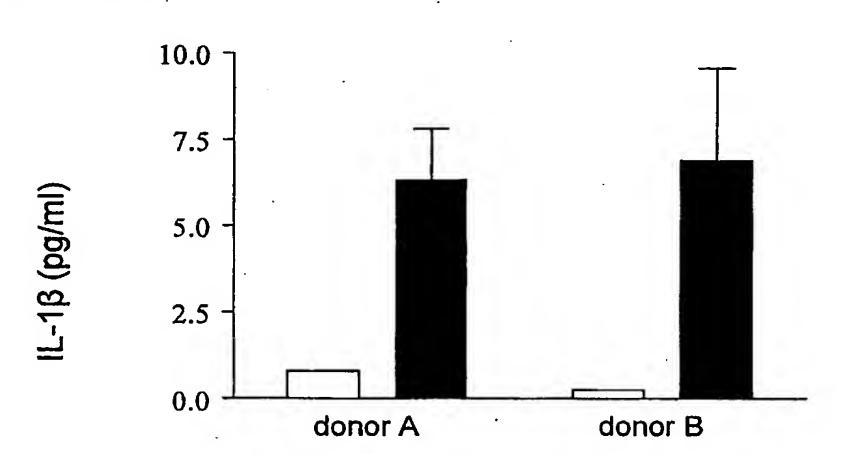


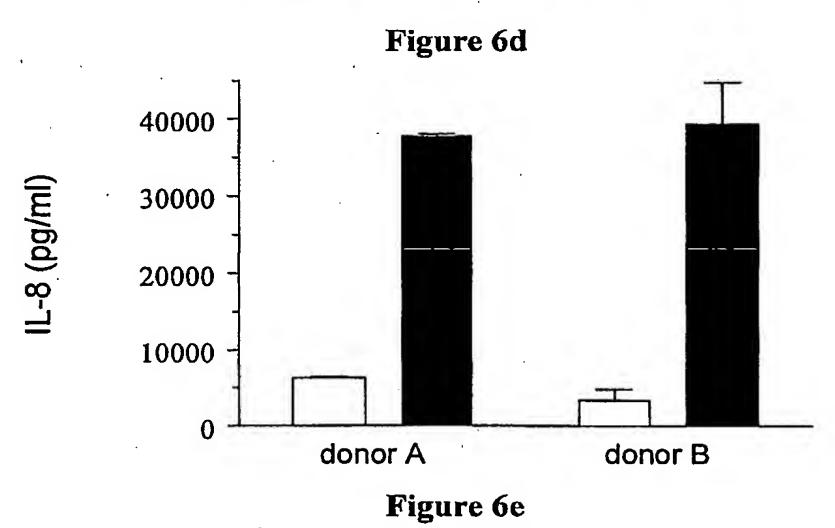
Figure 6

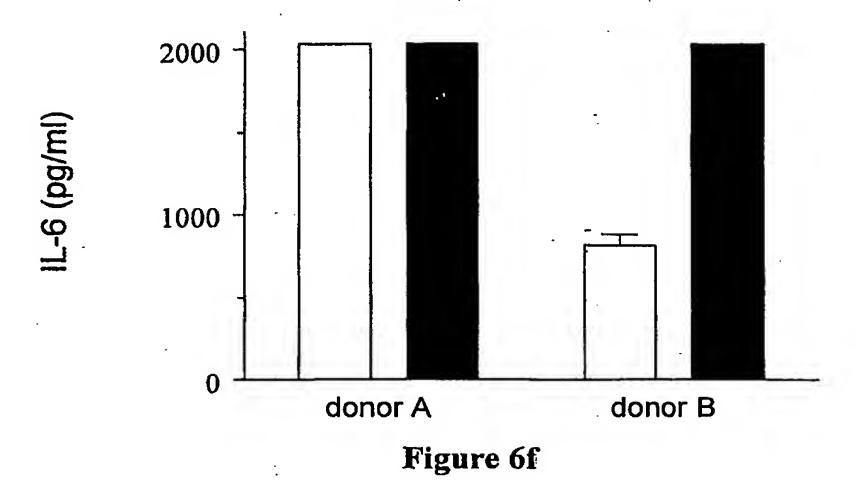
Release of chemokines and cytokines from single donor human peritoneal macrophages incubated with media control (□) or with PMAA-Na at 500 µg/ml (■) for 36 h











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Figure 7

Release of MIP-1β from human monocyte derived macrophages by endotoxin free PMAA-Na after 36 h from a single donor A

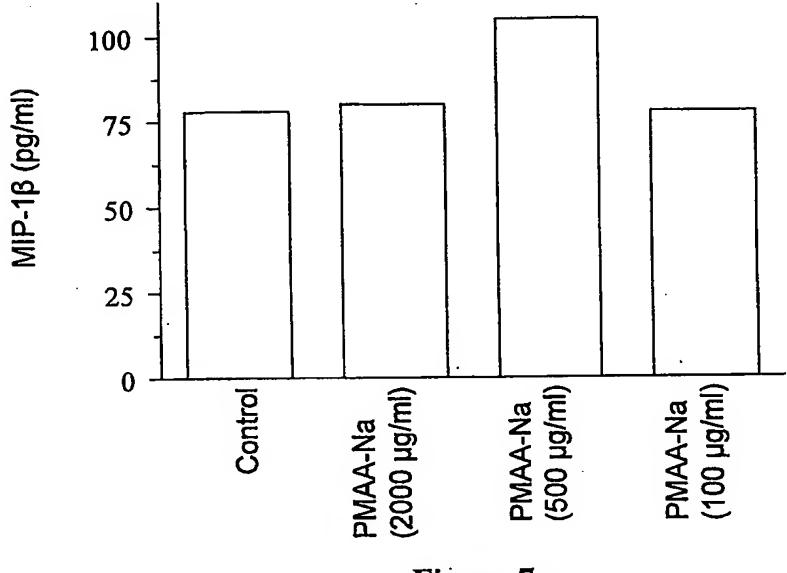


Figure 7a

Release of MIP-1β from human monocyte derived macrophages by endotoxin free PMAA-Na after 36 h from a single donor B

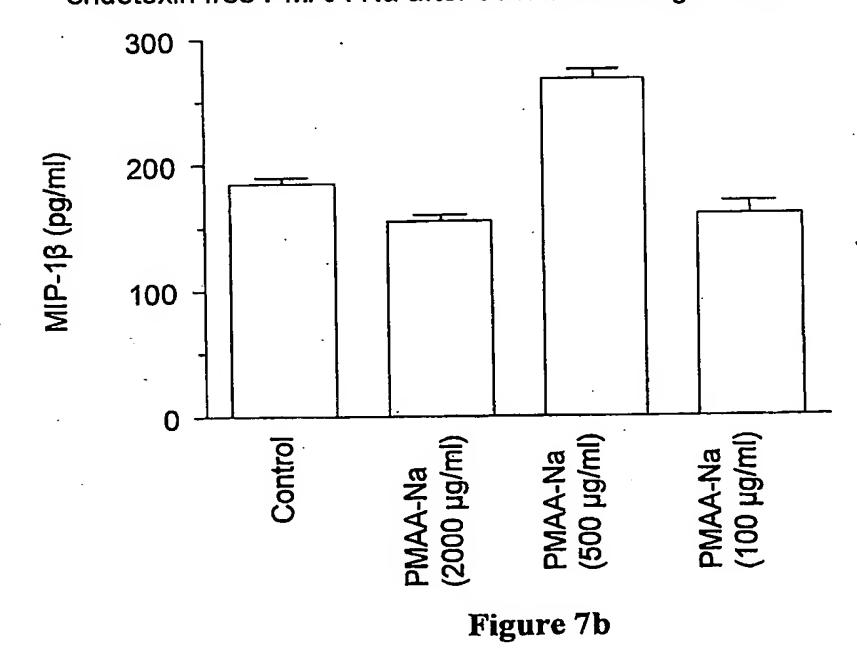


Figure 7 cont.

Release of MIP-1β from human monocyte derived macrophages by endotoxin free PMAA-Na after 36 h from a single donor C

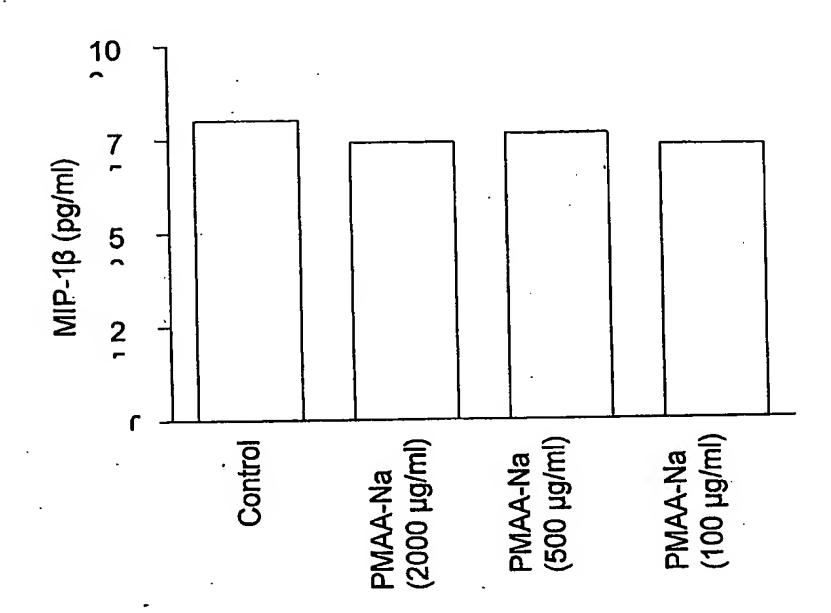
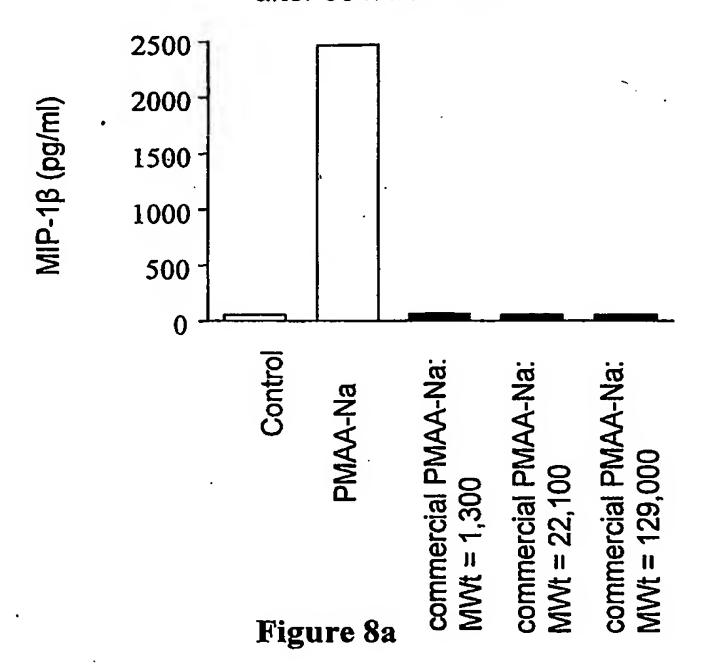


Figure 7c

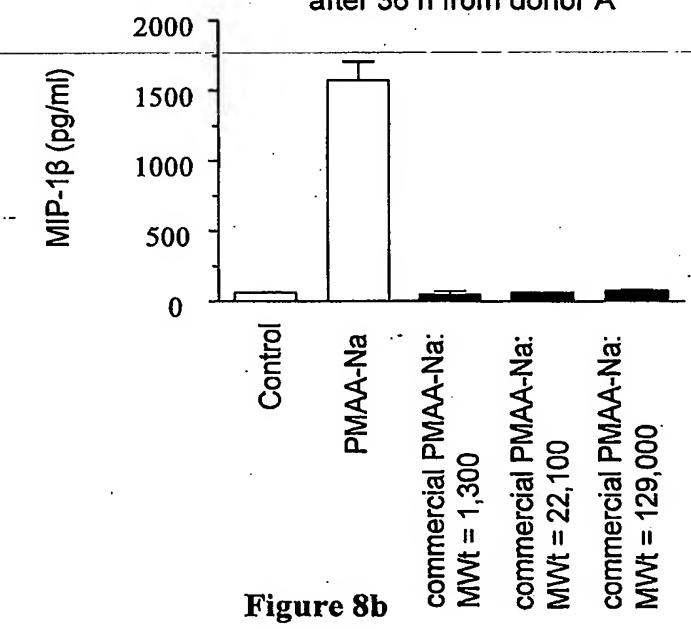
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Figure 8

Release of MIP-1β from human peritoneal macrophages by PMAA-Na (500 μg/ml) but not by commercially available PM AA-Na (500 μg/ml) after 36 h from donor A



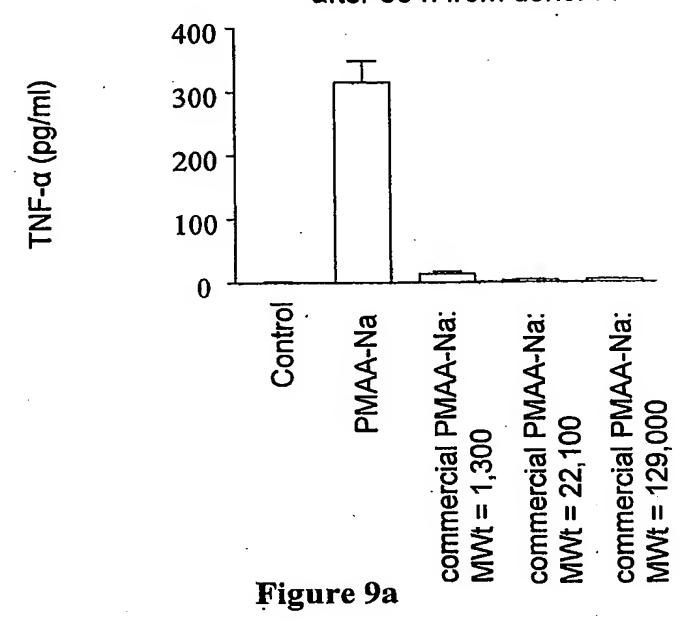
Release of MIP-1β from human peritoneal macrophages by PMAA-Na (500 μg/ml) but not by commercially available PM AA-Na (500 μg/ml) after 36 h from donor A



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Figure 9

Release of TNF-α from human peritoneal macrophages by PMAA-Na (500 μg/ml) but not by commercially available PM AA-Na (500 μg/ml) after 36 h from donor A



Release of TNF-α from human peritoneal macrophages by PMAA-Na (500 μg/ml) but not by commercially available PM AA-Na (500 μg/ml) after 36 h from donor A

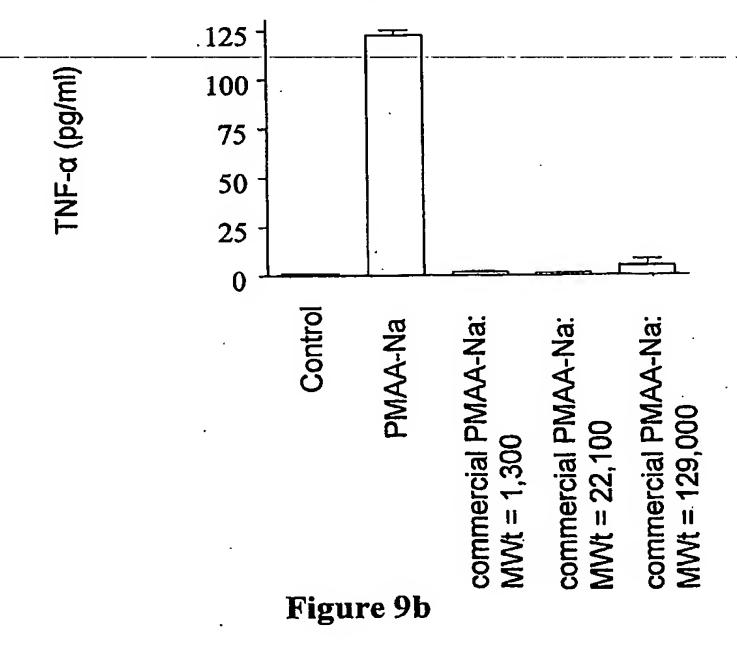


Figure 10



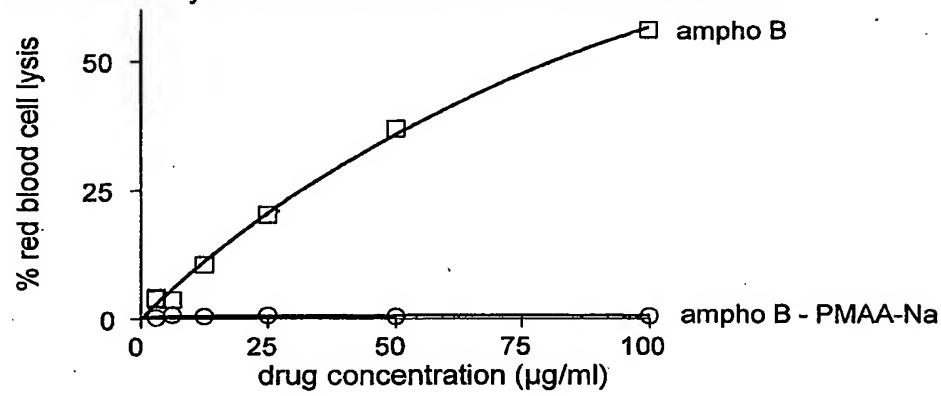


Figure 10a



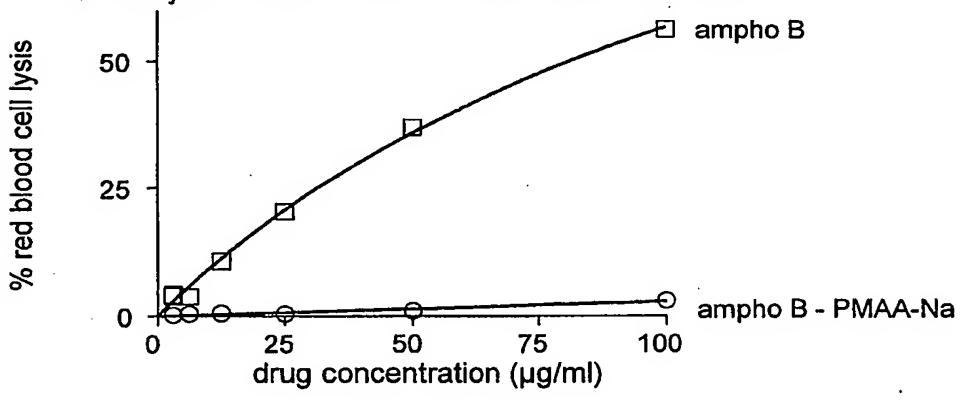


Figure 10b

Red blood cell lysis in donor C after a 1 hour incubation in RPMI

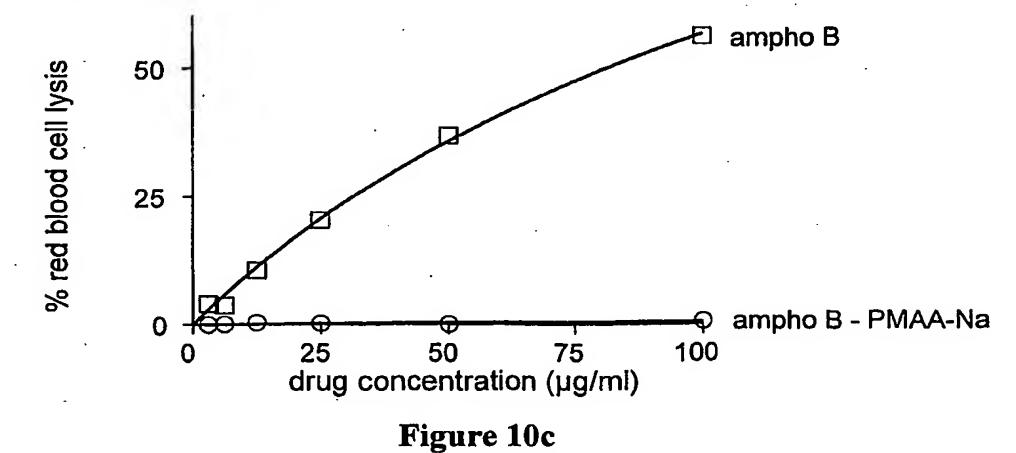
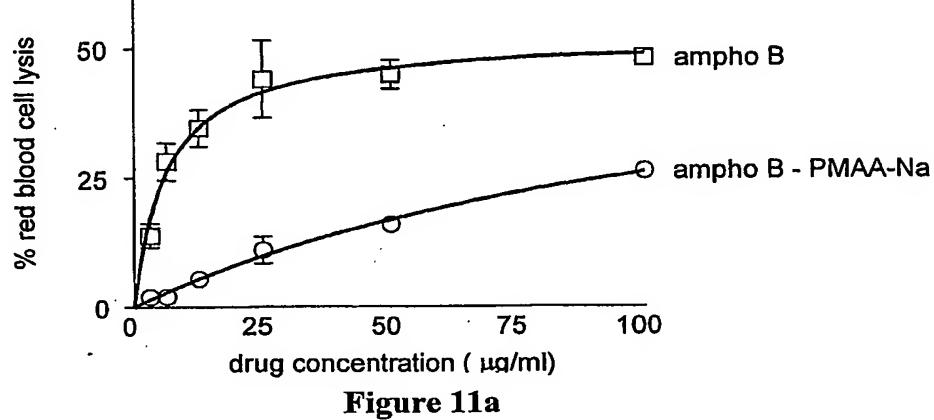


Figure 11

Red blood cell lysis in donor A after a 6 hour incubation in RPMI



Red blood cell lysis in donor B after a 6 hour incubation in RPMI

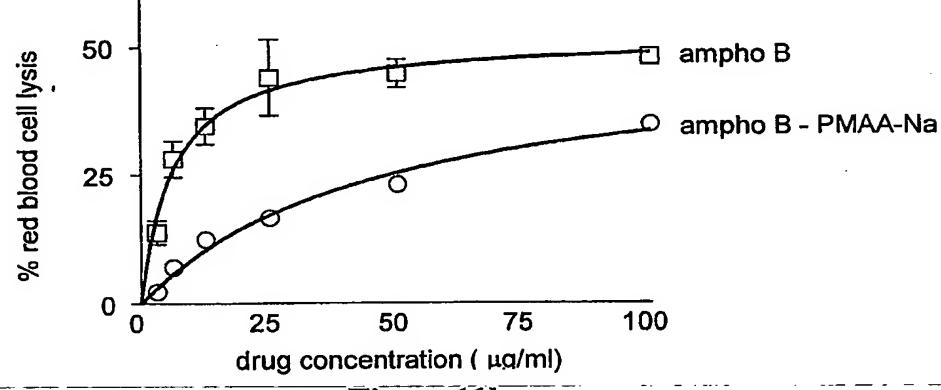


Figure 11b

Red cell lysis in donor C after a 6 hour incubation in RPMI

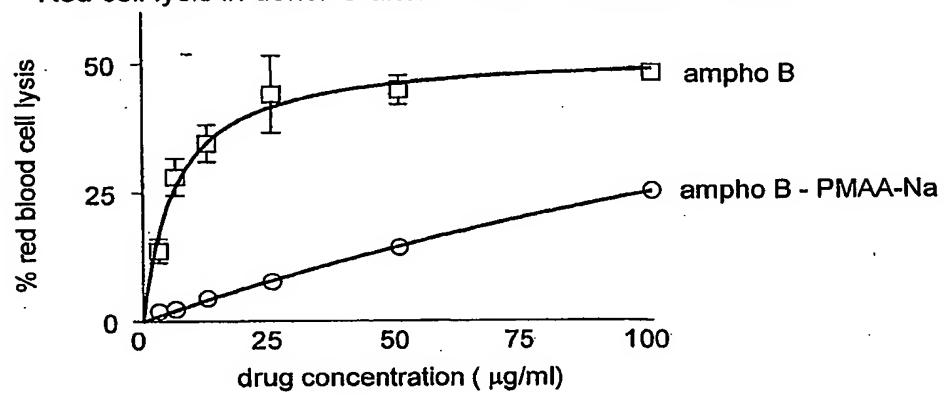


Figure 11c

Figure 12

Red blood cell lysis in donor A after a 24 hour incubation in RPMI

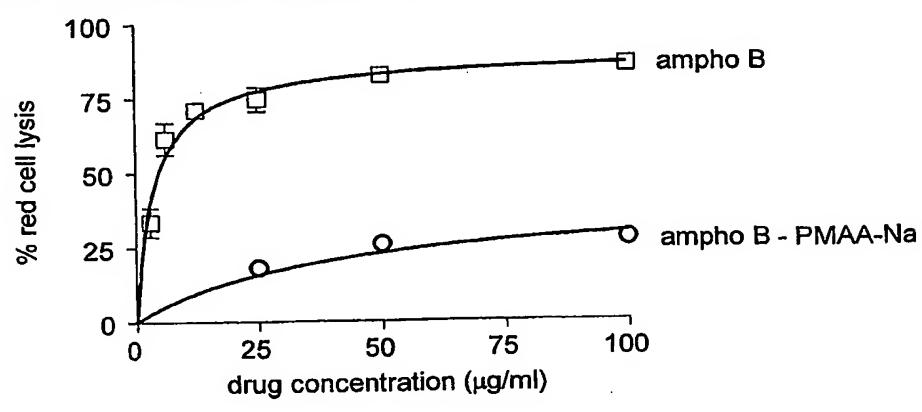


Figure 12a

Red cell lysis in donor C after a 24 hour incubation in RPMI

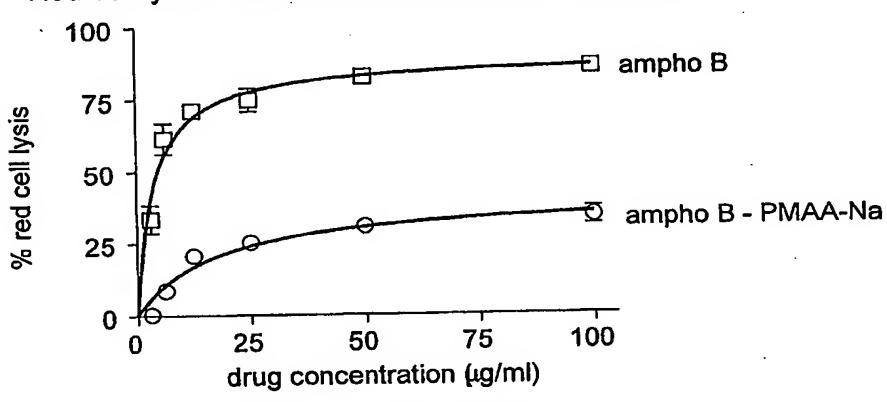


Figure 12b

Red blood cell lysis in donor C after a 24 hour incubation in RPMI

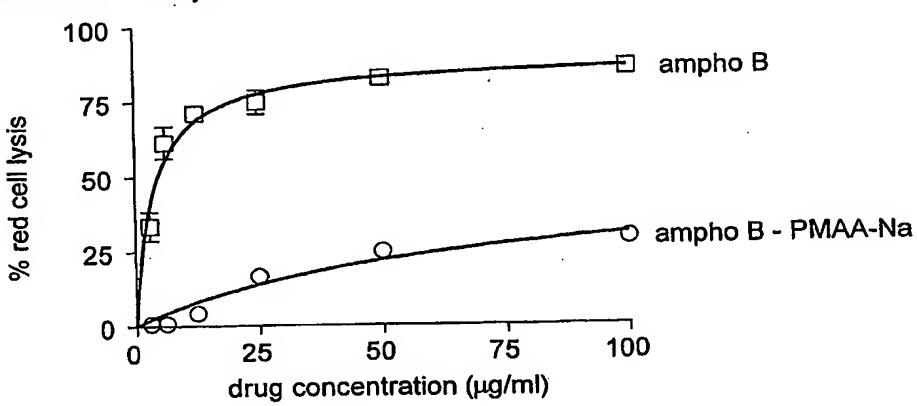


Figure 12c

Figure 13

Red blood cell lysis in single donor after incubation with the amphotericin B - PMAA-Na preparation in RPMI

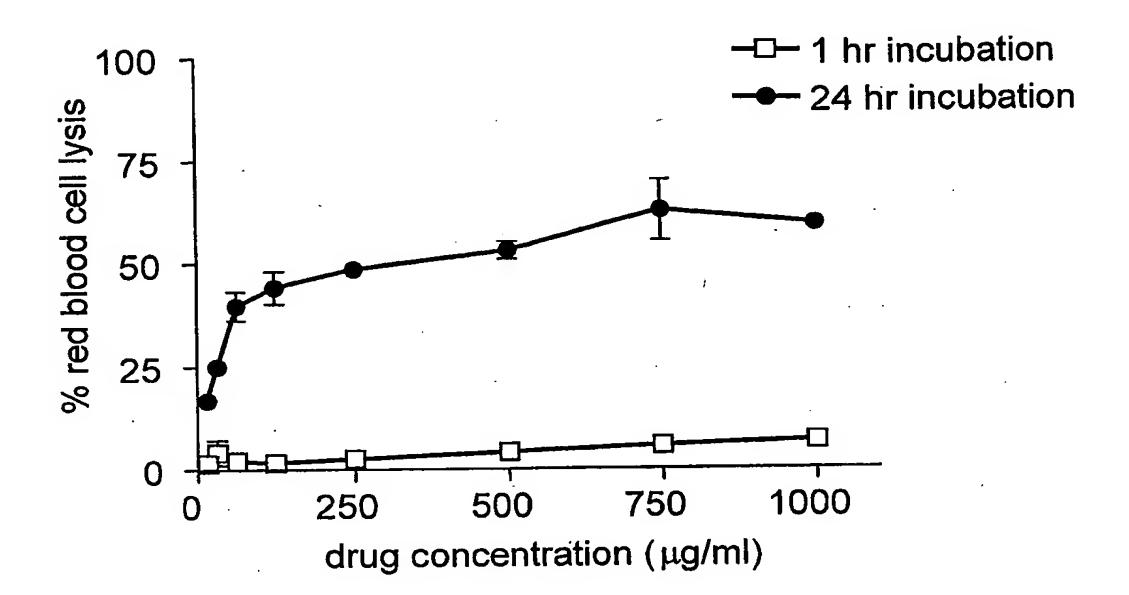
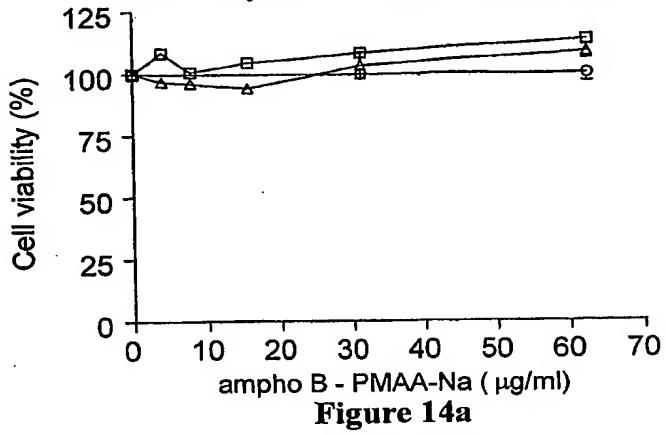
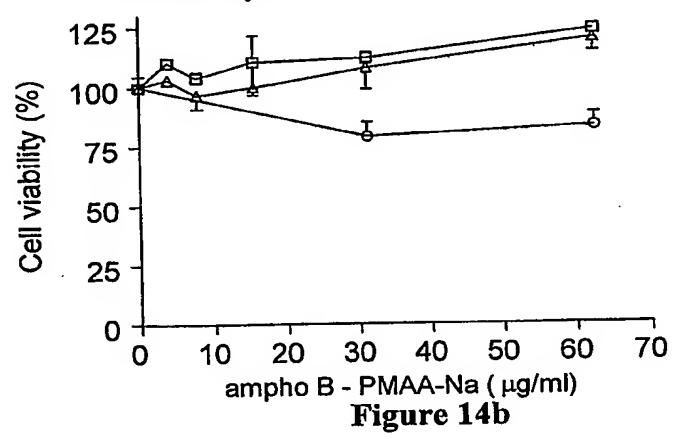


Figure 14

Lack of toxicity of the amphotericin B - PMAA-Na preparation after 1 day of culture with PBMN cells (n=3)



Lack of toxicity of the amphotericin B - PMAA-Na preparation after 2 days of culture with PBMN cells (n=3)



Lack of the toxicity of amphotericin B - PMAA-Na preparation after 6 days of culture with PBMN cells (n=2)

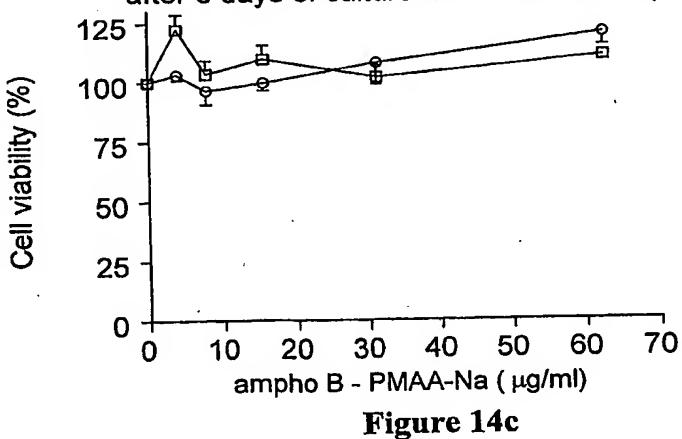


Figure 15

Toxicity of the amphotericin B - PMAA-Na preparation after 2 days of culture with monocyte derived macrophages

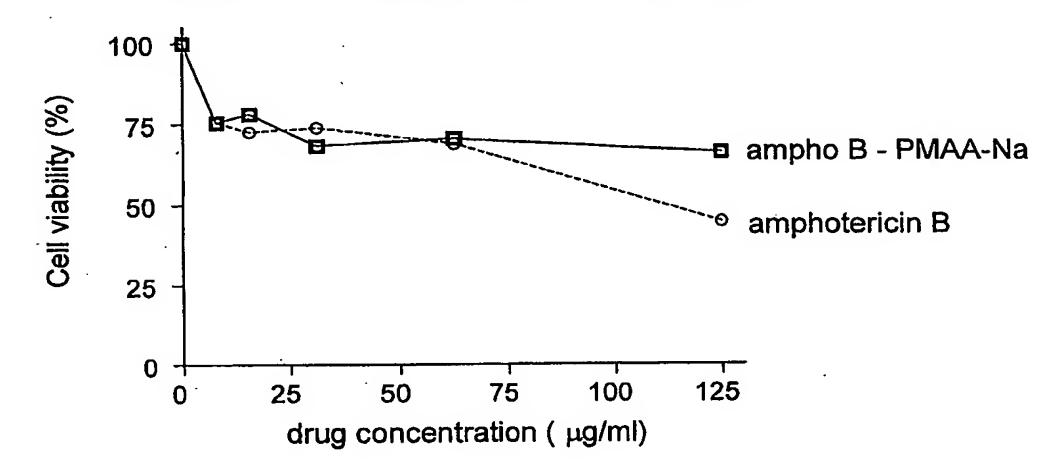
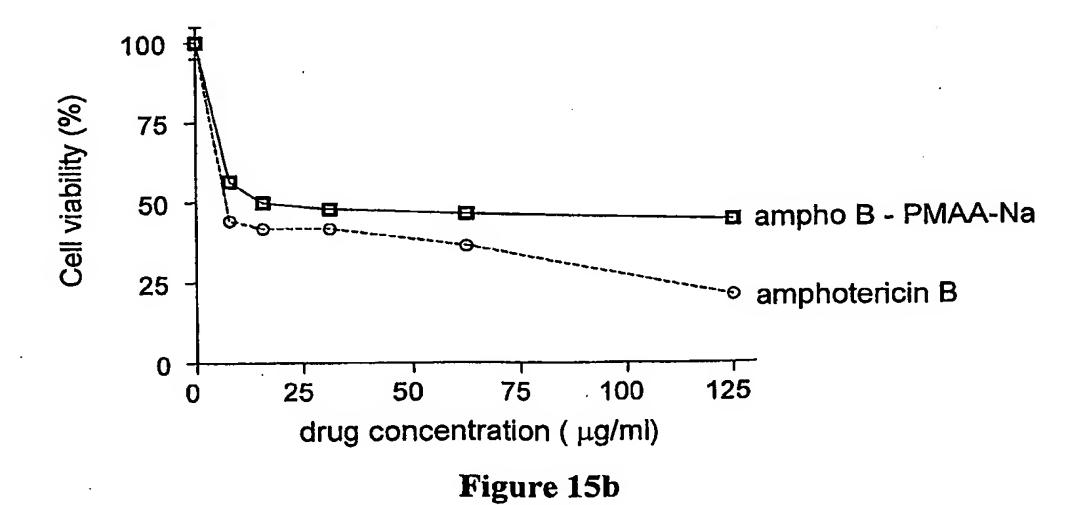


Figure 15a

Toxicity of the amphotericin B - PMAA-Na preparation after 3 days of culture with monocyte derived macrophages

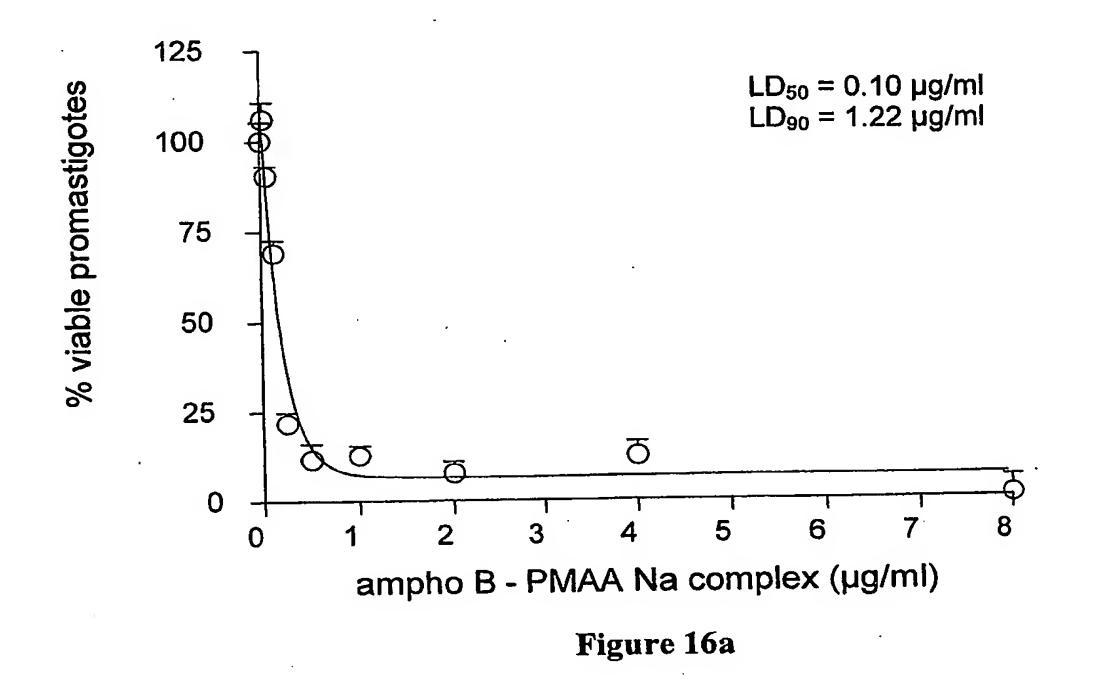


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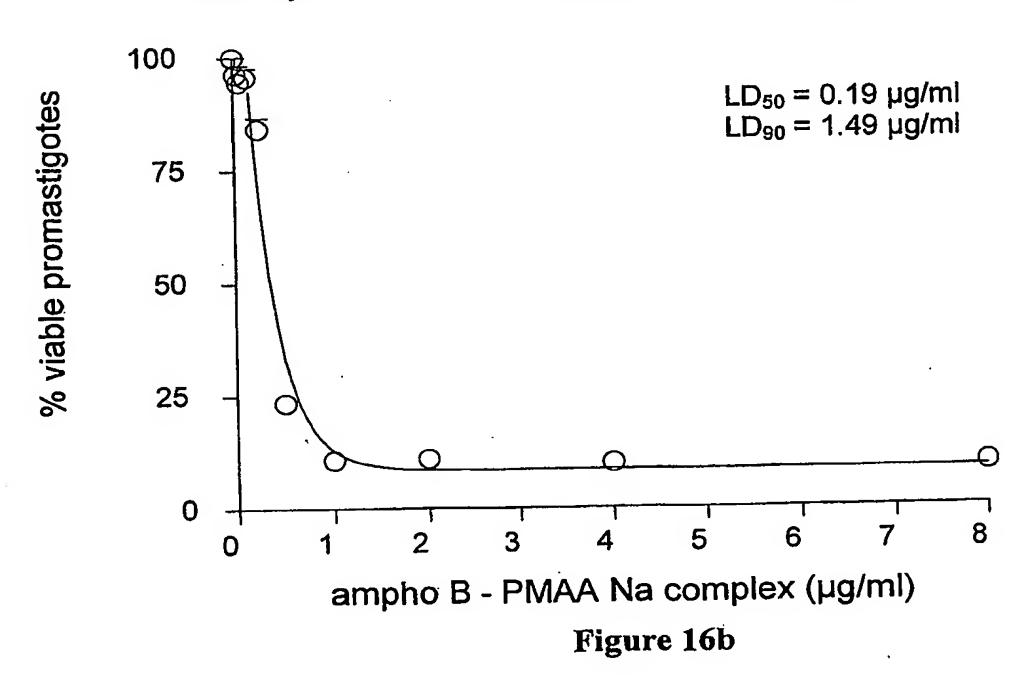
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Figure 16

Viability of Leishmania mexicana promastigotes after 2 days



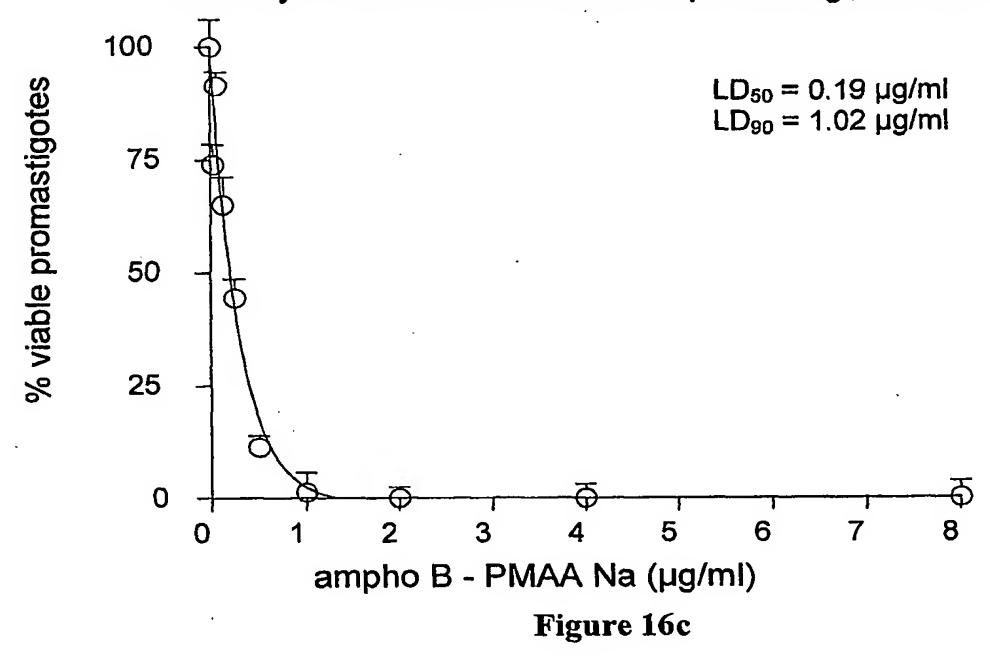
Viability of Leishmania mexicana promastigotes after 2 days



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Figure 16 cont.

Viability of Leishmania mexicana promastigotes after 2 days



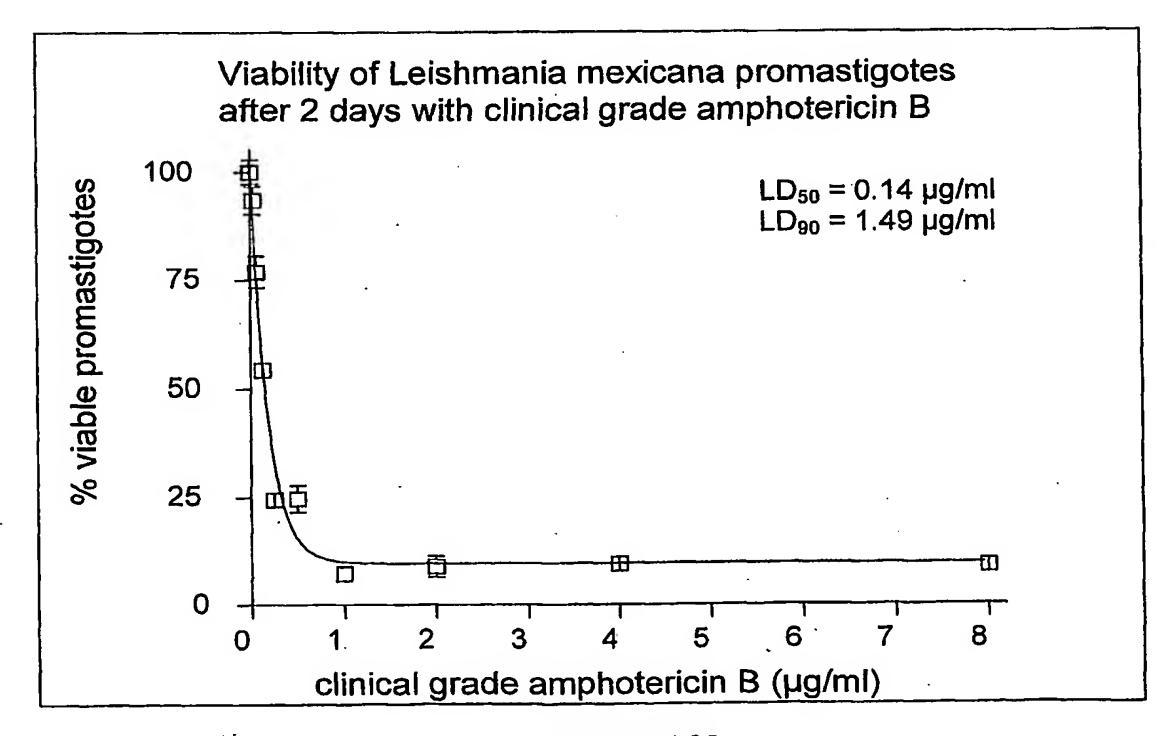


Figure 16d

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Figure 17
Viability of Leishmania donovani promasigotes after 2 days

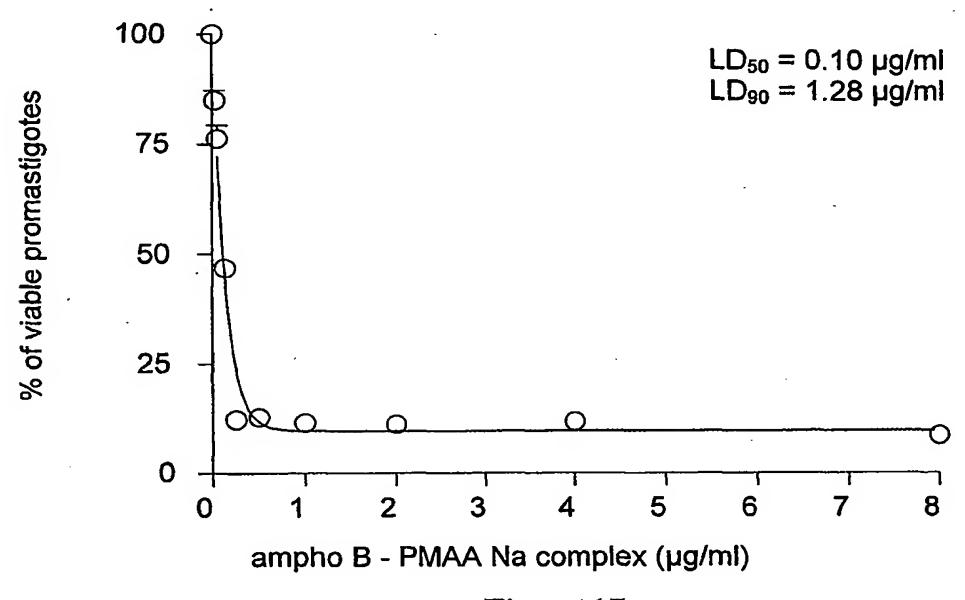


Figure 17a

Viability of Leishmania donovani promasigotes after 2 days

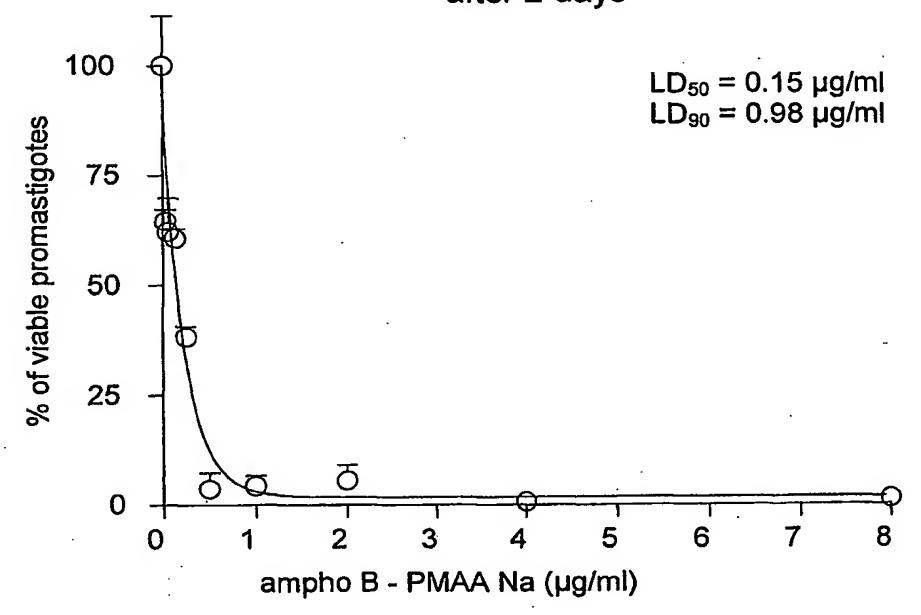
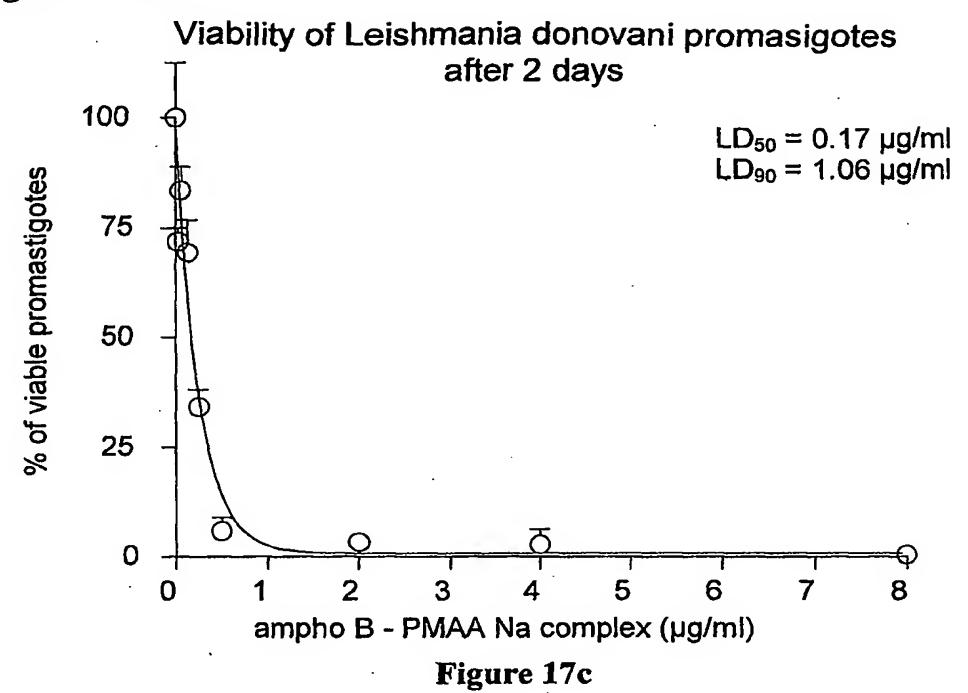


Figure 17b

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Figure 17 cont.



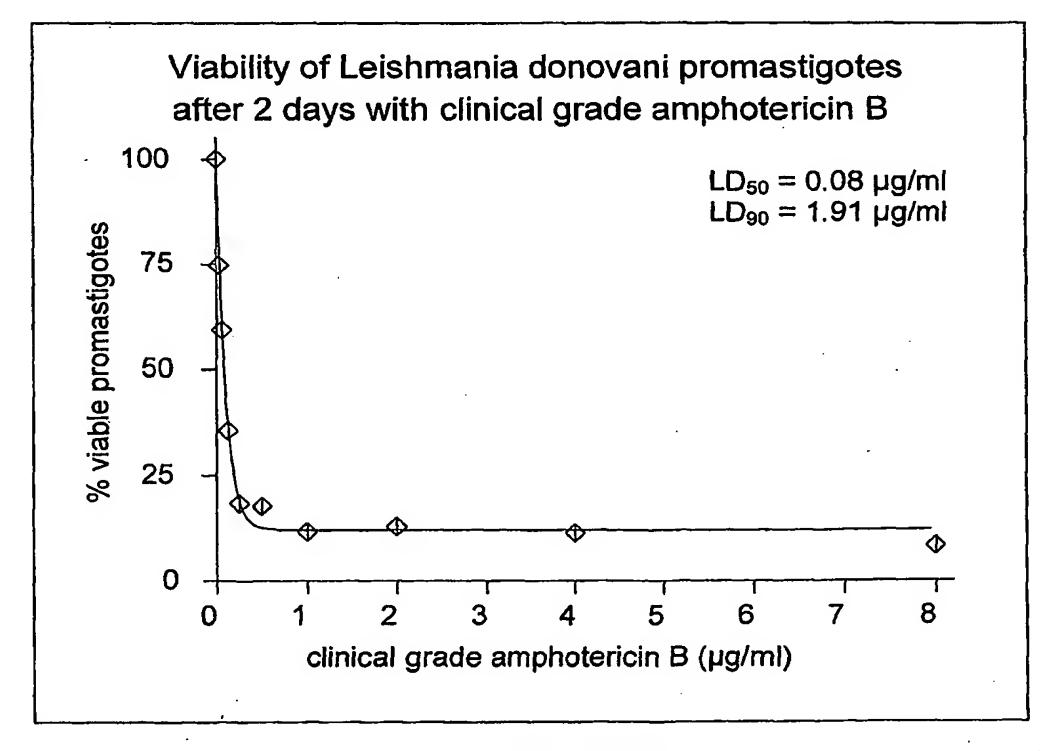
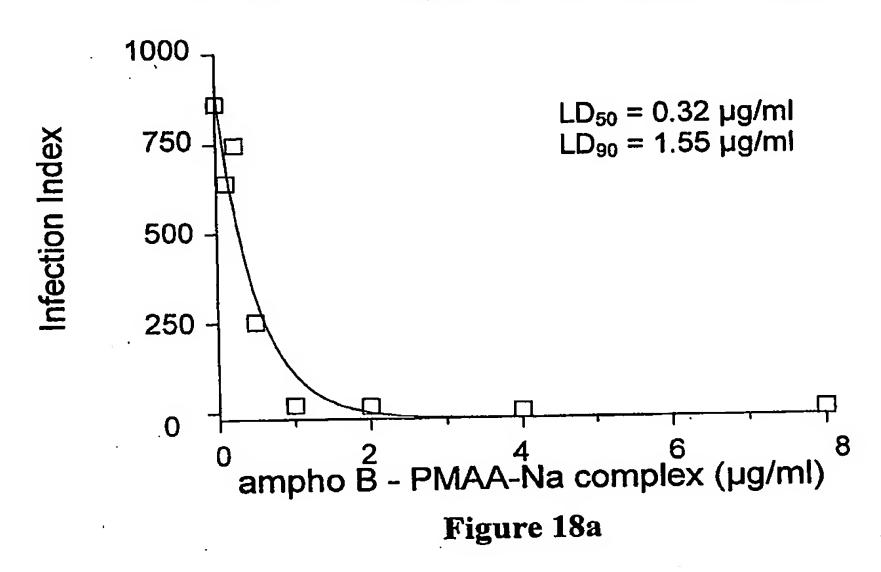


Figure 17d

Figure 18

Inhibition of intracellular Leishmania mexicana amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages



Inhibition of intracellular Leishmania mexicana amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

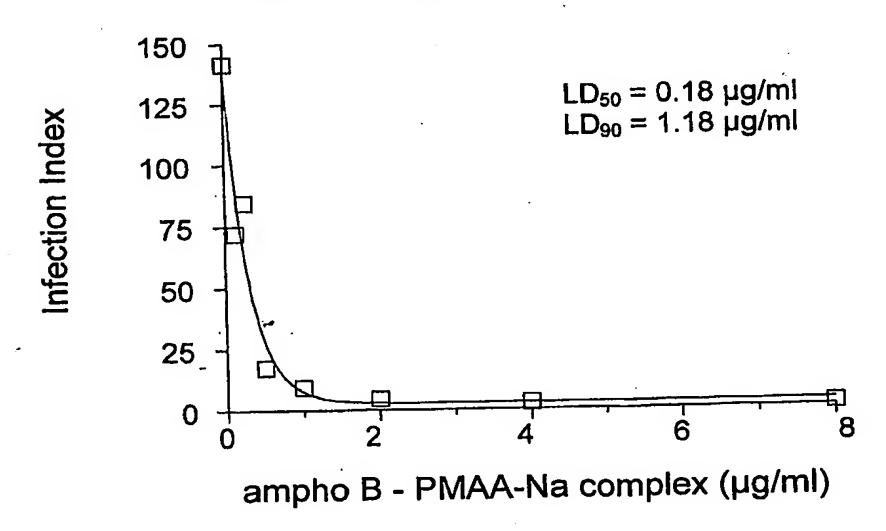


Figure 18b

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Figure 18 cont.

Inhibition of intracellular Leishmania mexicana amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

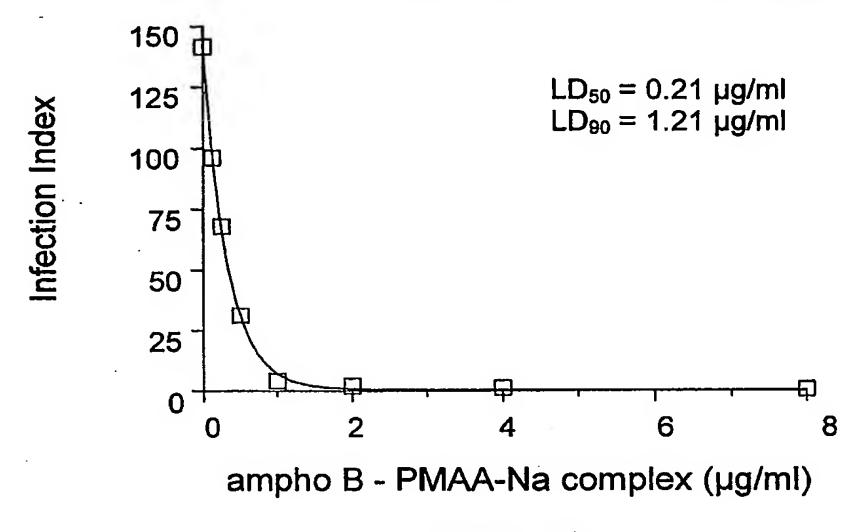


Figure 18c

Inhibition of intracellular Leishmania mexicana amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

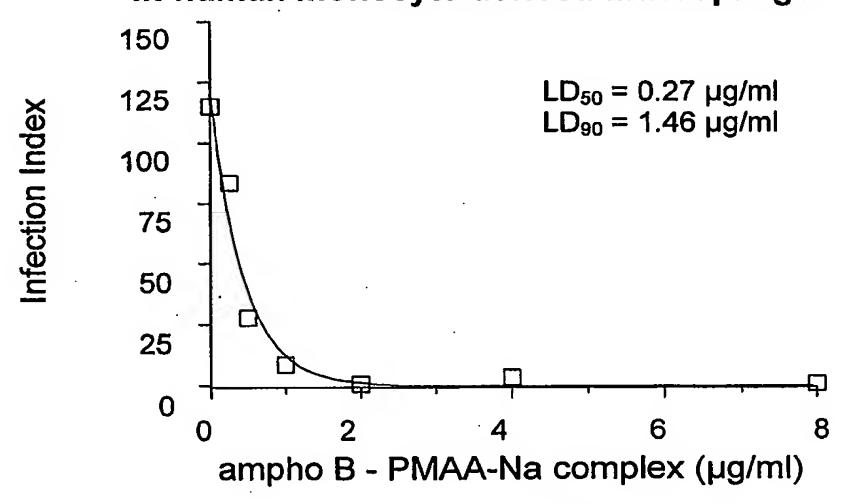
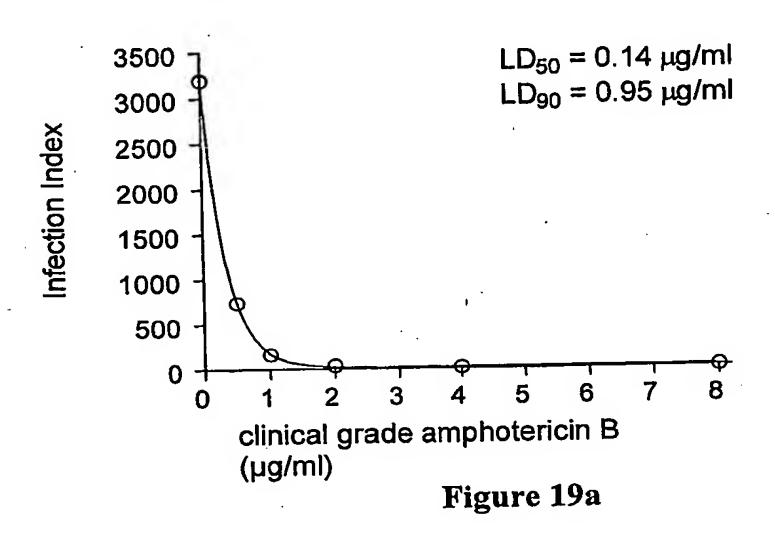


Figure 18d

Figure 19

Inhibition of intracellular Leishmania mexicana amastigote growth by clincal grade amphotericin B in human monocyte derived macrophages



Inhibition of intracellular Leishmania mexicana amastigote growth by Ambisome (Gilead Sciences) in human monocyte derived macrophages

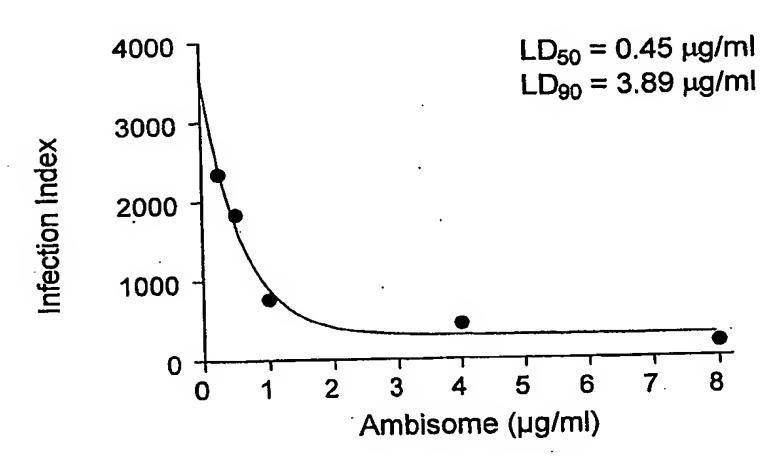
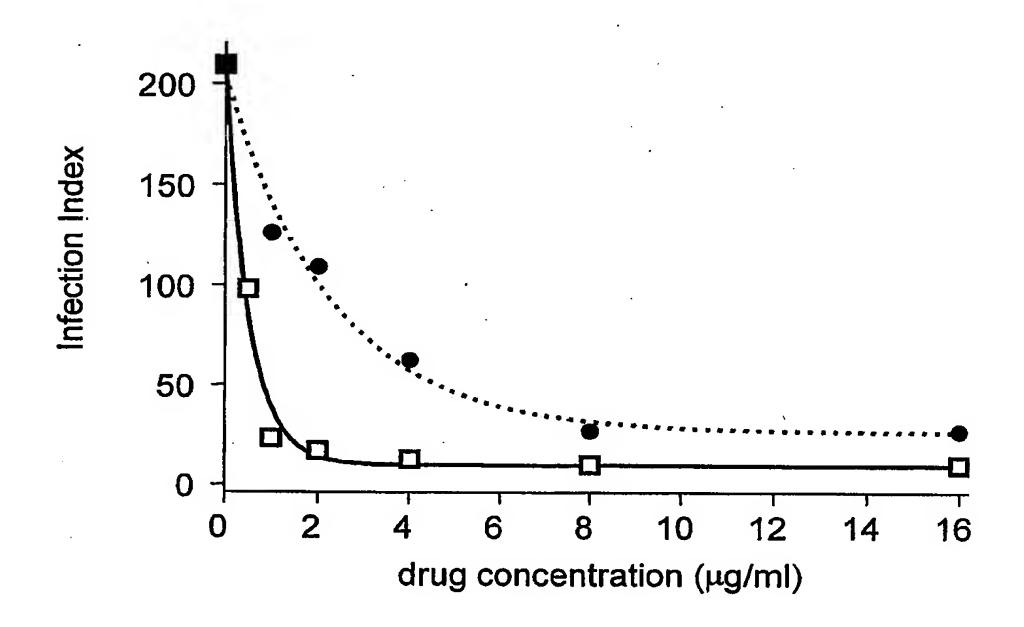


Figure 19b

Figure 20

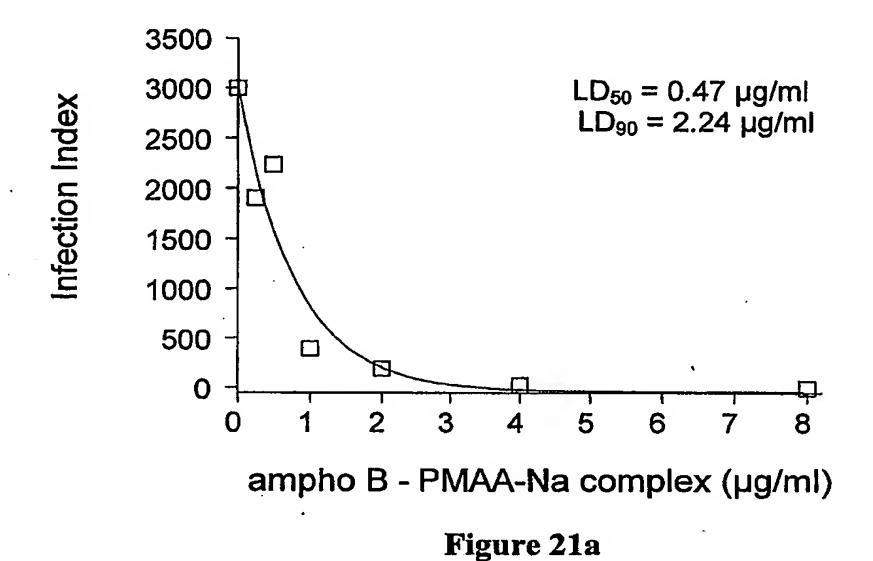
Inhibition of intracellular Leishmania mexicana amastigote growth in human macrophages by amphotericin B - PMAA-Na compared to AmBisome



```
— amphotericin B - PMAA-Na = 0.3 μg/ml IC<sub>50</sub> = 0.7 μg/ml ·······Ambisome (Gilead Sciences): IC<sub>50</sub> = 1.7 μg/ml
```

Figure 21

Inhibition of intracellular Leishmania donovani amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages



Inhibition of intracellular Leishmania donovani amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

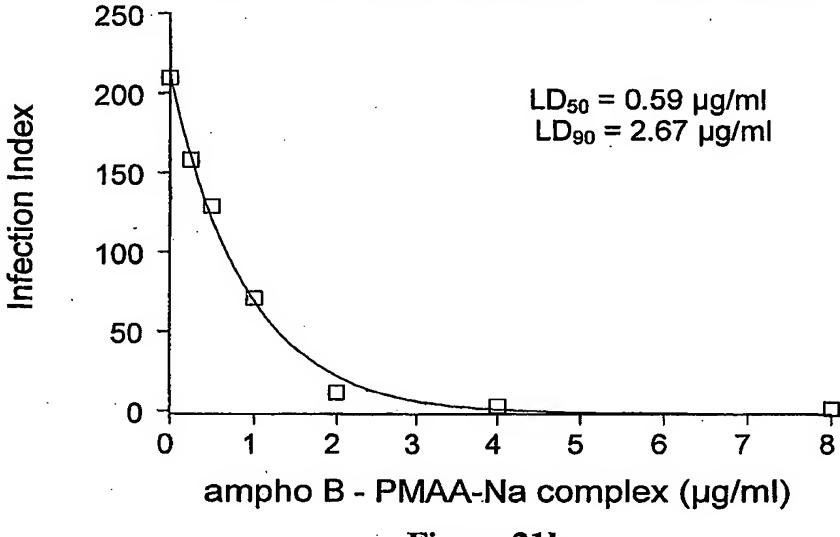


Figure 21b

Figure 21 cont.

Inhibition of intracellular Leishmania donovani amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

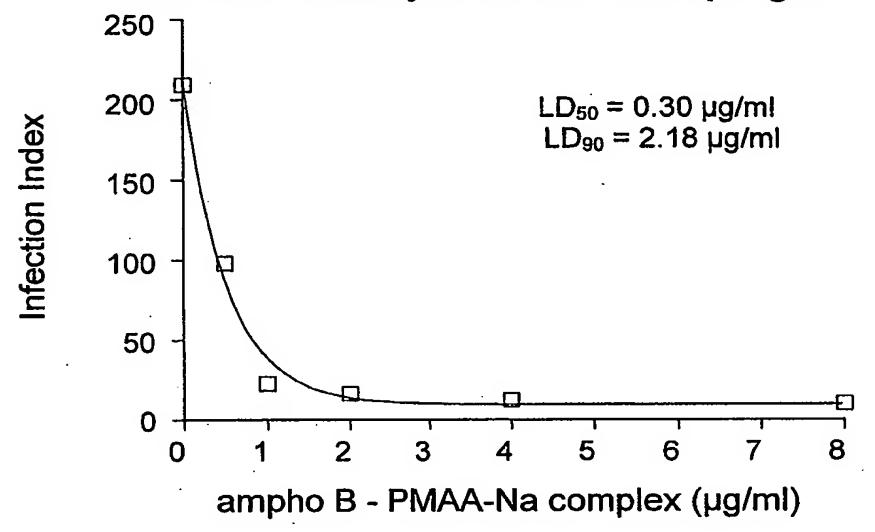


Figure 21c

Inhibition of intracellular Leishmania donovani amastigote growth by the amphotericin B-PMAA-Na preparation in human monocyte derived macrophages

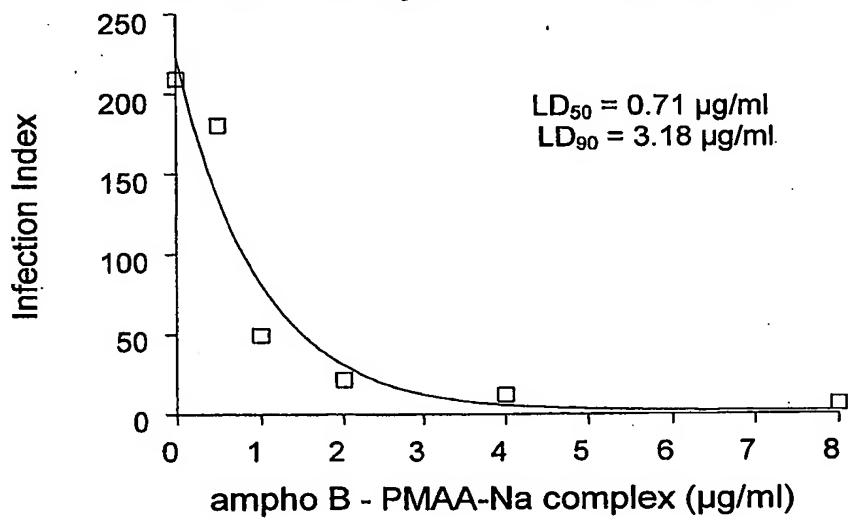


Figure 21d

Figure 22

Inhibition of intracellular Leishmania donovani amastigote growth by clinical grade amphotericin B in human monocyte derived macrophages

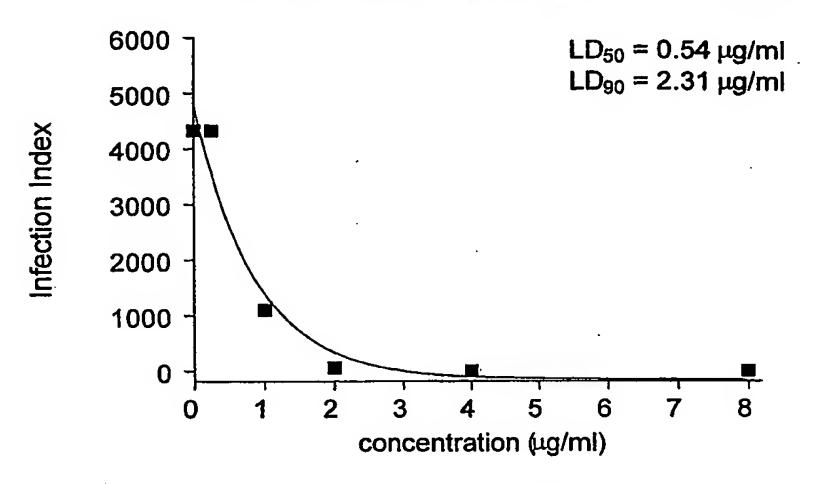


Figure 22a

Inhibition of intracellular Leishmania donovani amastigote growth by Ambisome (Gilead Sciences) in human monocyte derived macrophages

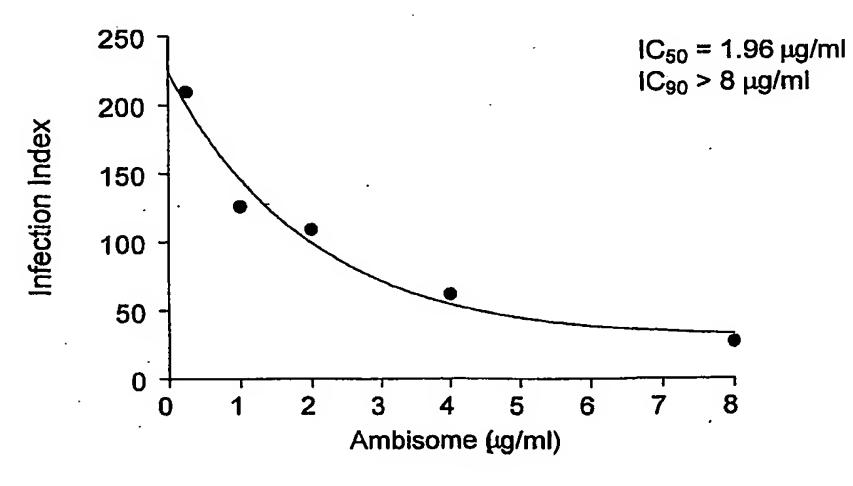
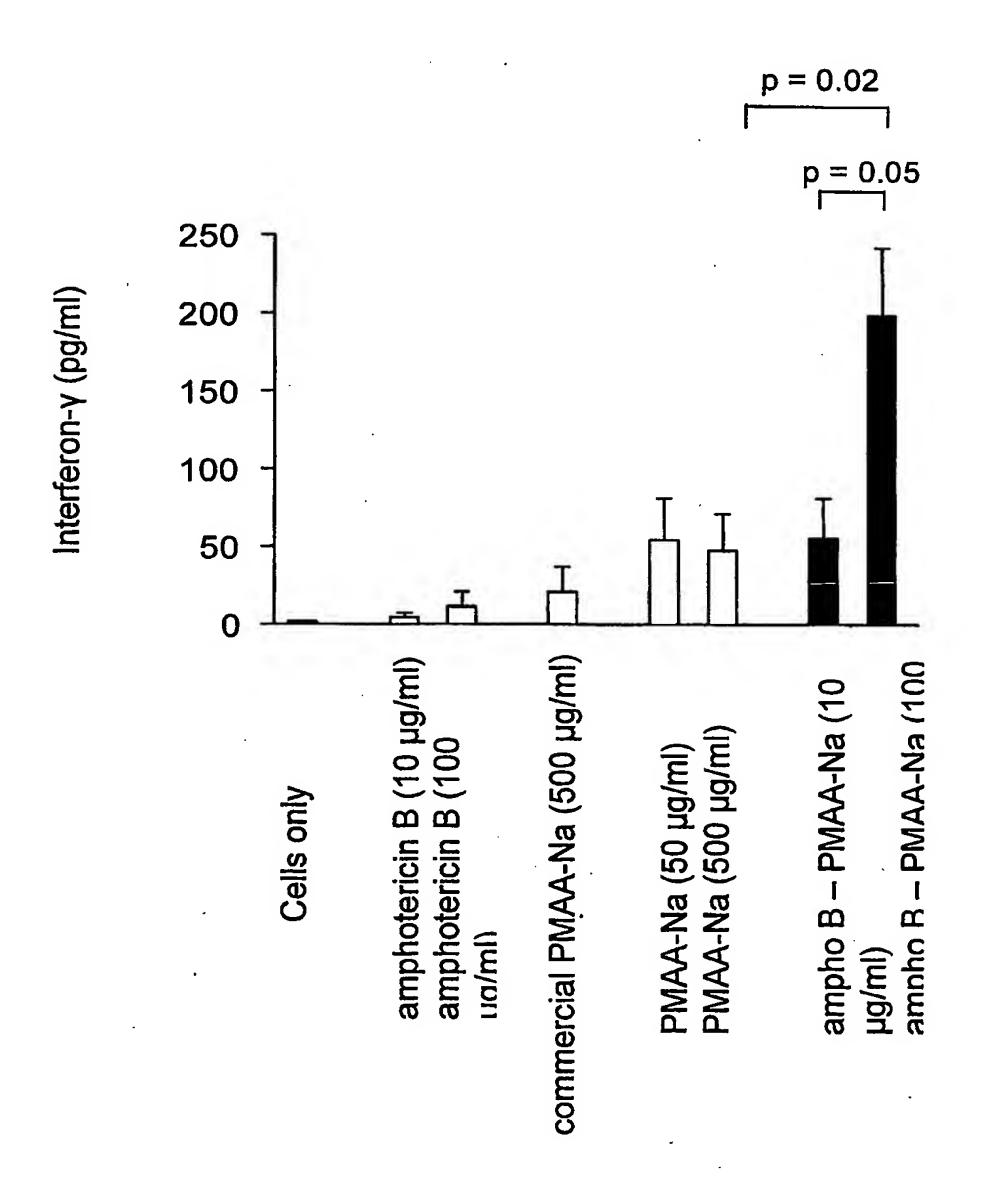


Figure 22b

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Figure 23

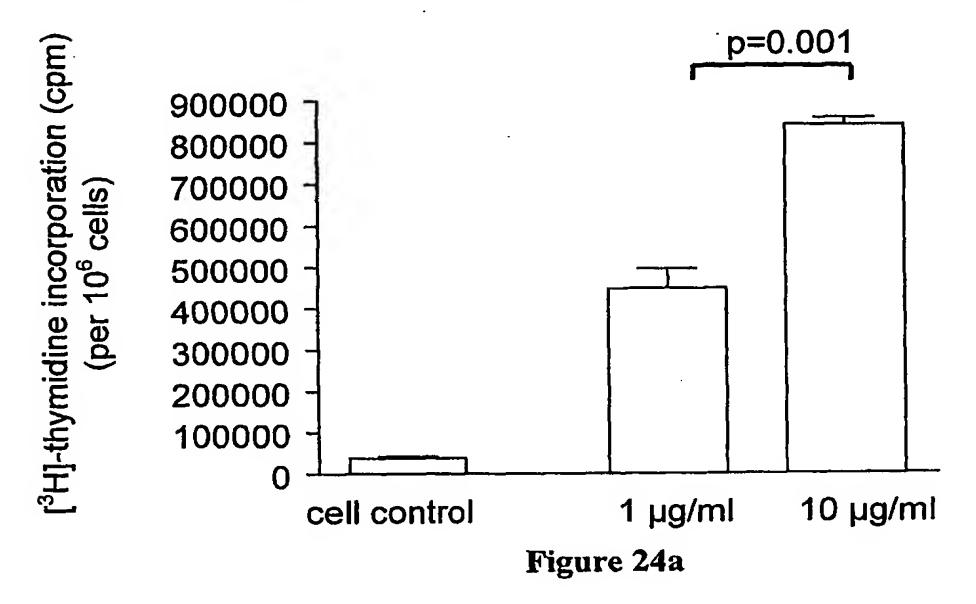
Release of interferon- γ from human peritoneal macrophags after culture with different compounds for 24 h



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Figure 24

PBMC proliferation after 6 days incubation of the cells of donor A with the tuberculin PPD - PMAA-Na preparation



PBMC proliferation after 5 days stimulation with antigen of the cells from donor B

p=0.002

p=0.007

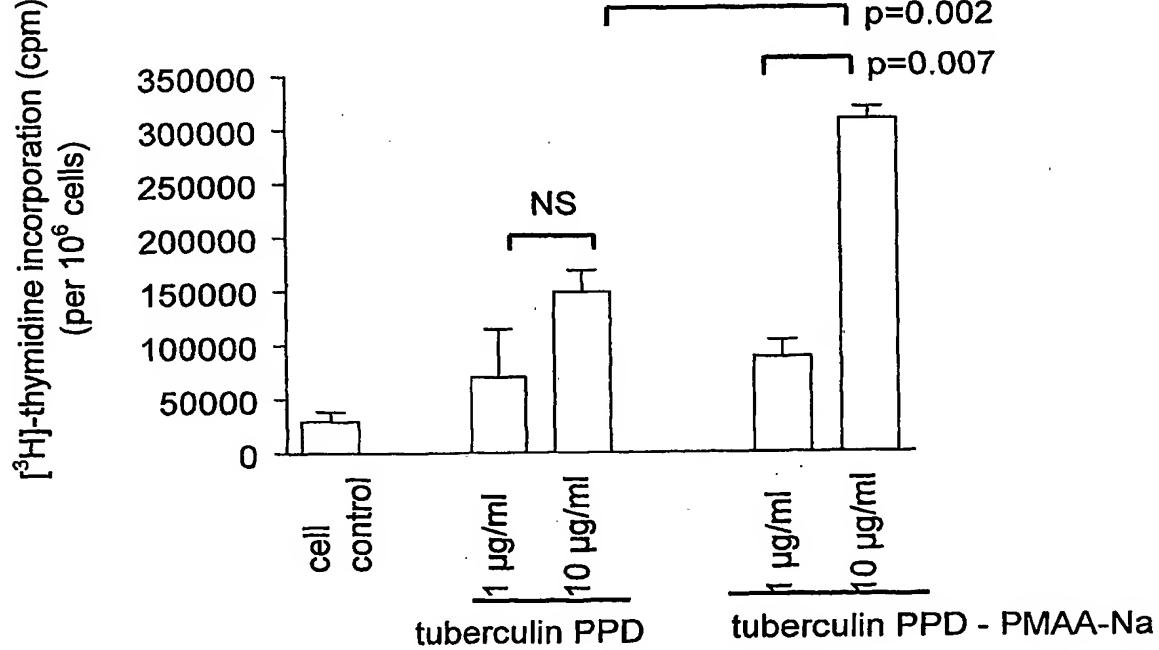
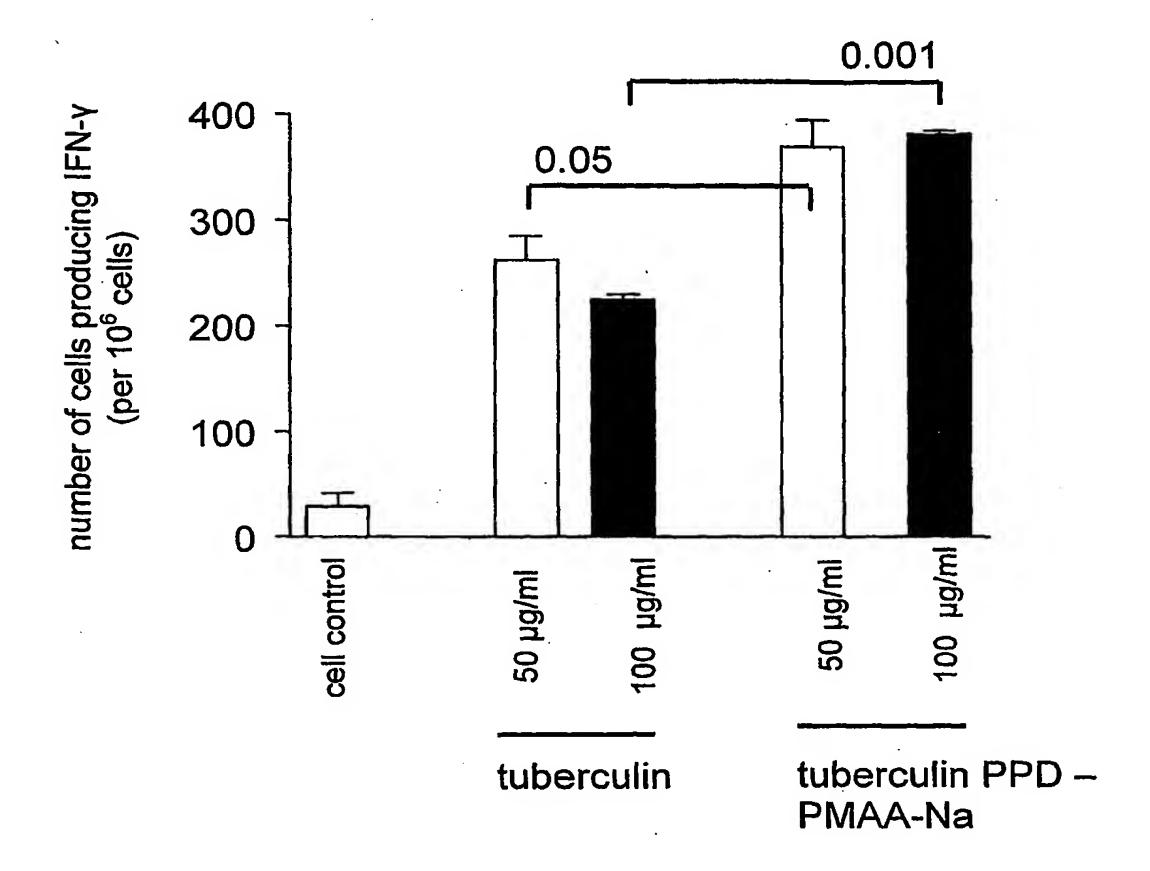


Figure 24b

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Figure 25

IFN-γ production by human PBMCs that were stimulated with antigen from donor A for 24 h



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Figure 26

Survival of L. donovani amastigotes in mouse liver macrophages after intravenous treatment with several different preparations of amphotericin B

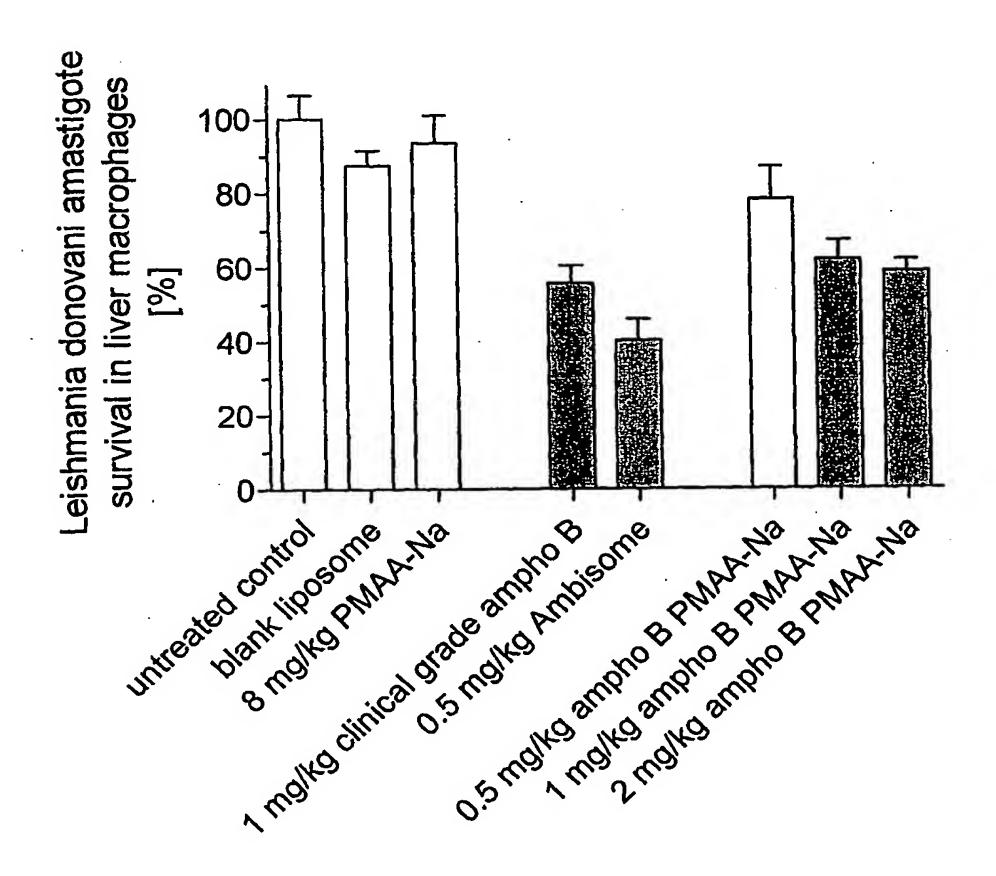


Figure 27

Lack of toxicity of PMAA-Na to single donor red blood cells (n=3)

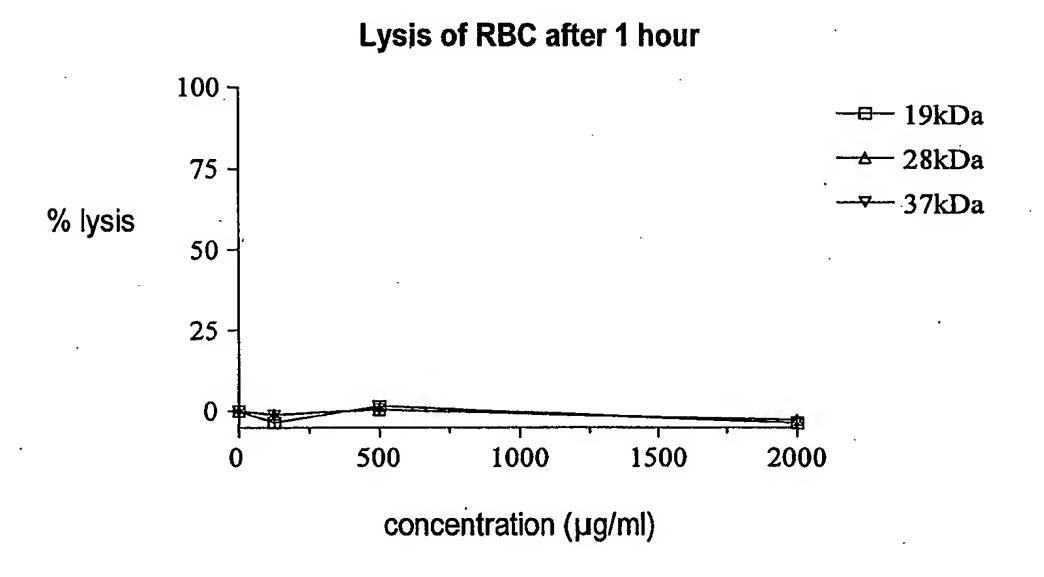


Figure 27a

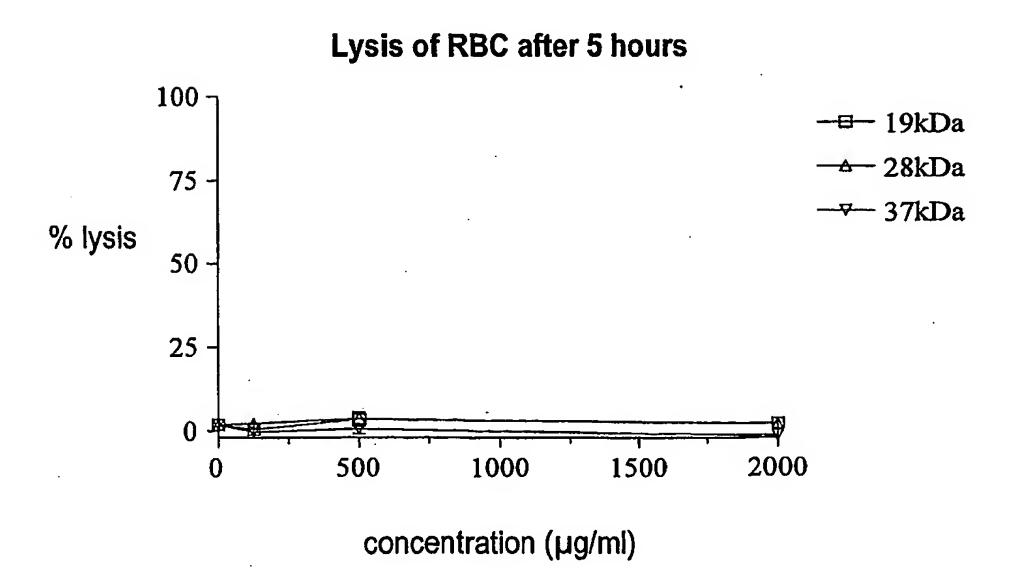


Figure 27b

Figure 27 cont.

Lysis of RBC after 24 hr incubation with sample

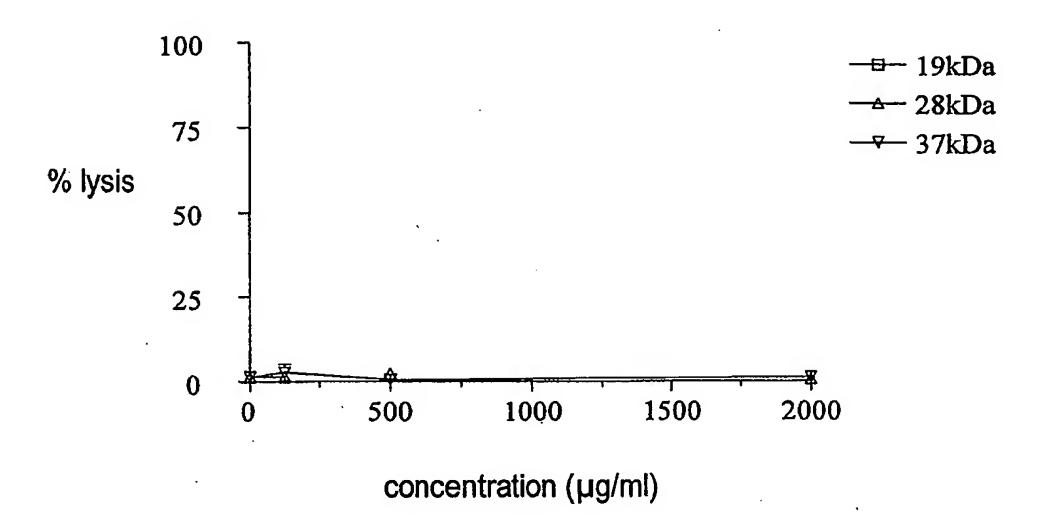


Figure 27c

Figure 28

Lack of toxicity of PMAA-NA to single donor PBMCs after a 1 day incubation (n=3)

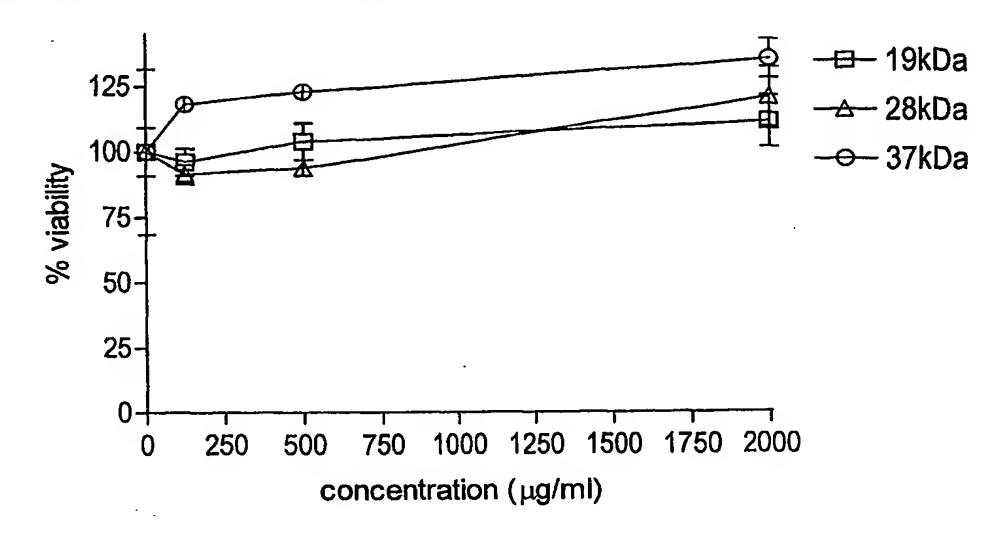


Figure 28a

Lack of toxicity of PMAA-NA to single donor PBMCs after a 2 day incubation (n=3)

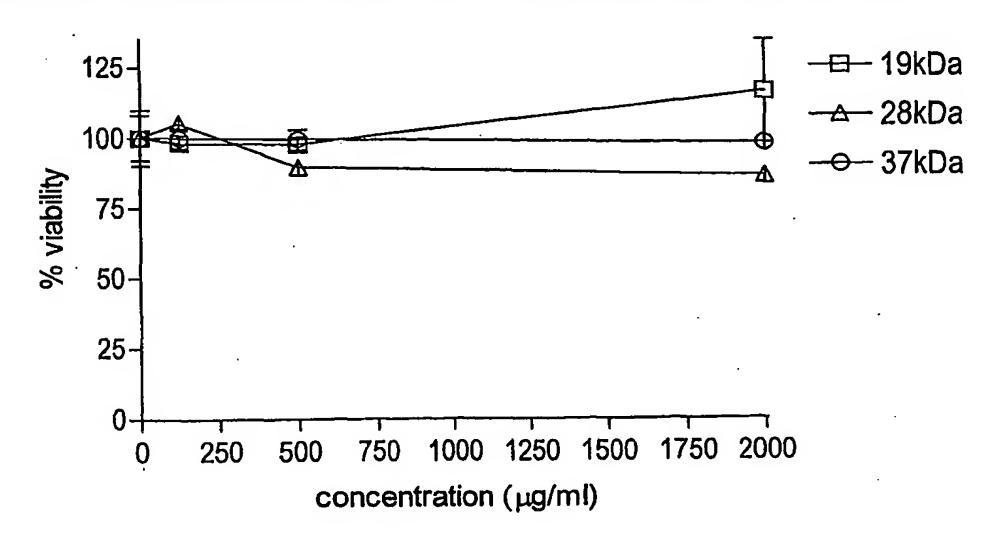


Figure 28b

Figure 29

Lysis of red blood cells after incubation with ampho B - PMAA-Na stored as a lyophilised powder at 4°C for 4 months (n=2)

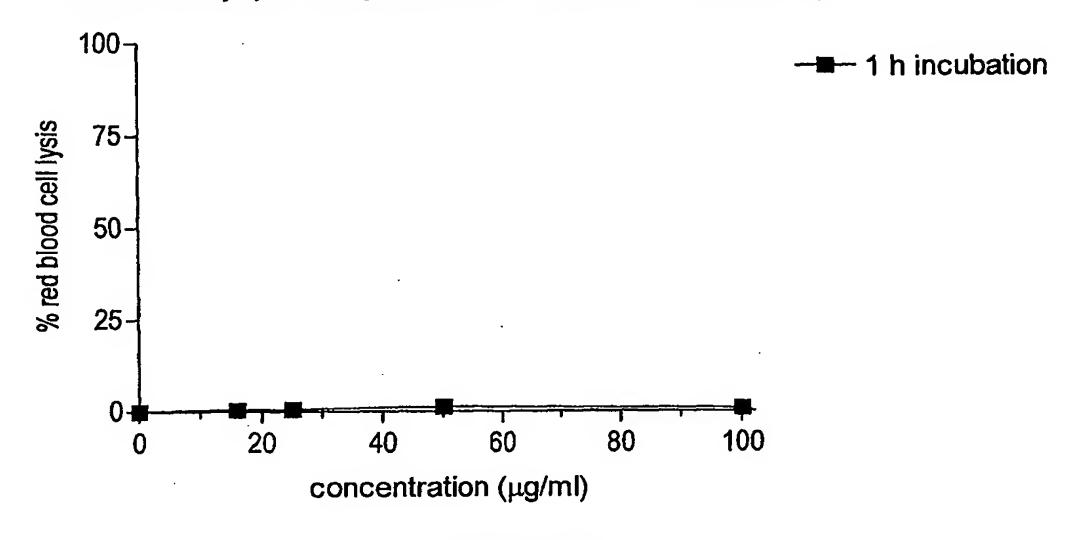


Figure 29a

Lysis of red blood cells after a 1 h incubation with amphotericin B - PMAA-Na stored in 5% dextrose at 4°C for 7 months

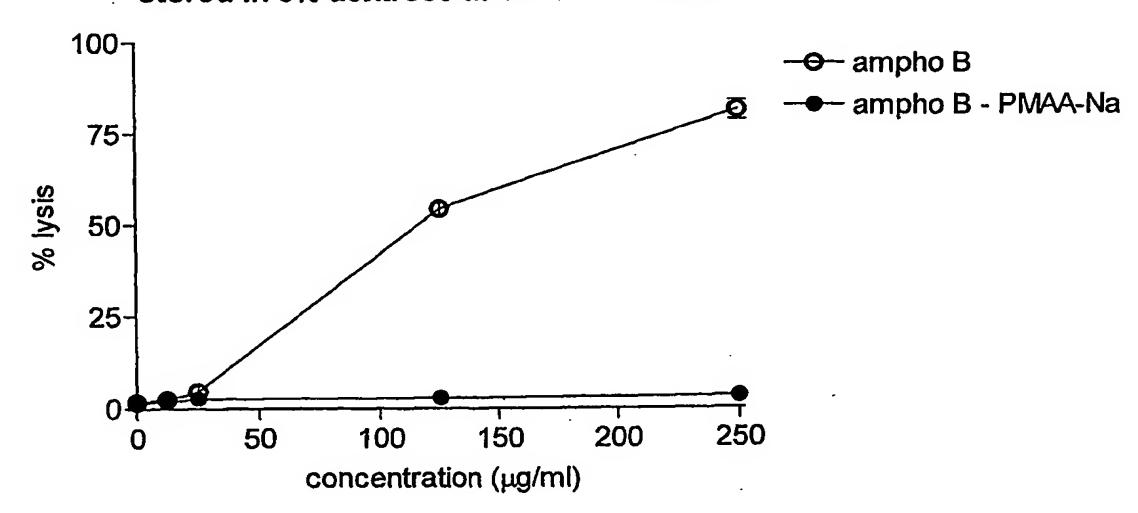


Figure 29b

Figure 30

Inhibition of L. mexicana amastigote growth in human MDMs after 3 d incubation with amphotericin B - PMAA-Na stored as a lyophilised powder at 4°C for 4 months

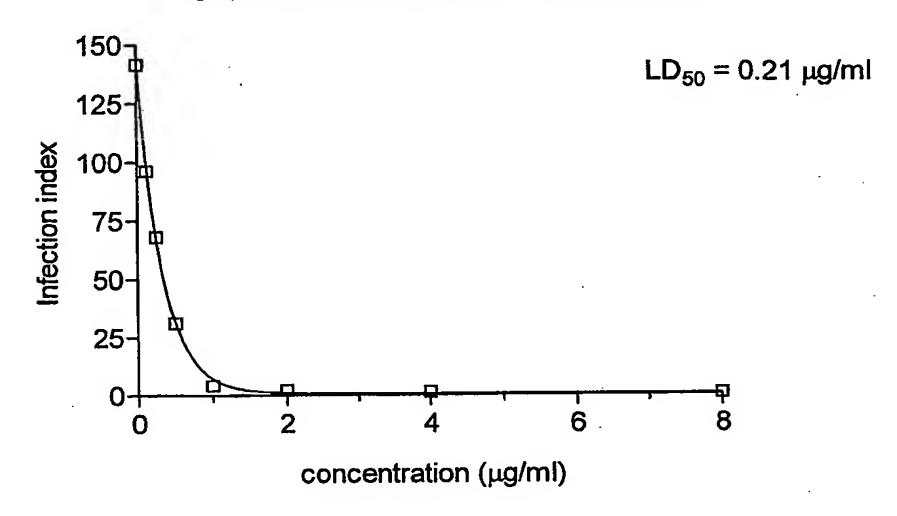


Figure 30a

Viability of *L.mexicana* promastigotes after 2 d incubation with amphotericin B - PMAA-Na stored in 5% dextrose at 4°C for 7 months

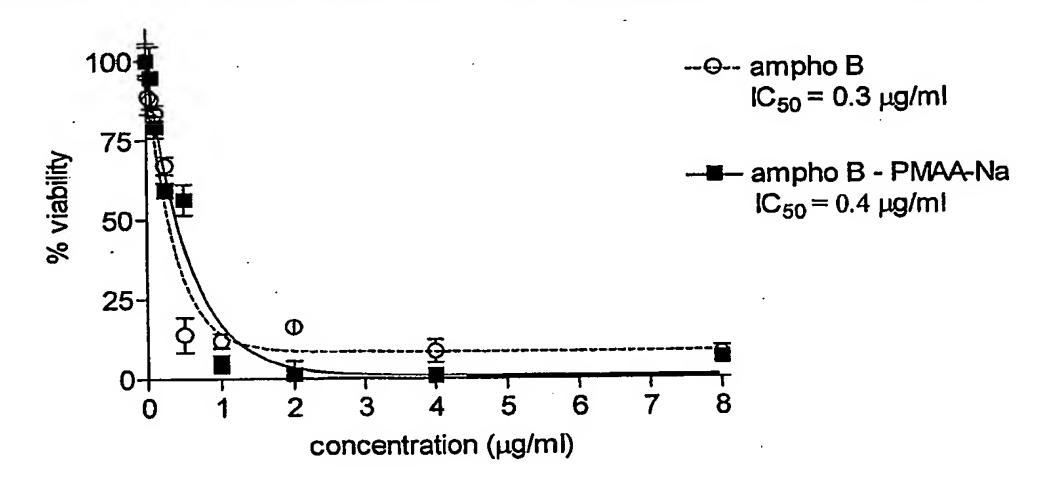


Figure 30b

Figure 31

Inhibition of *C. neoformans* var *neoformans* clinical isolate 1 in monocyte derived macrophages after 3 days

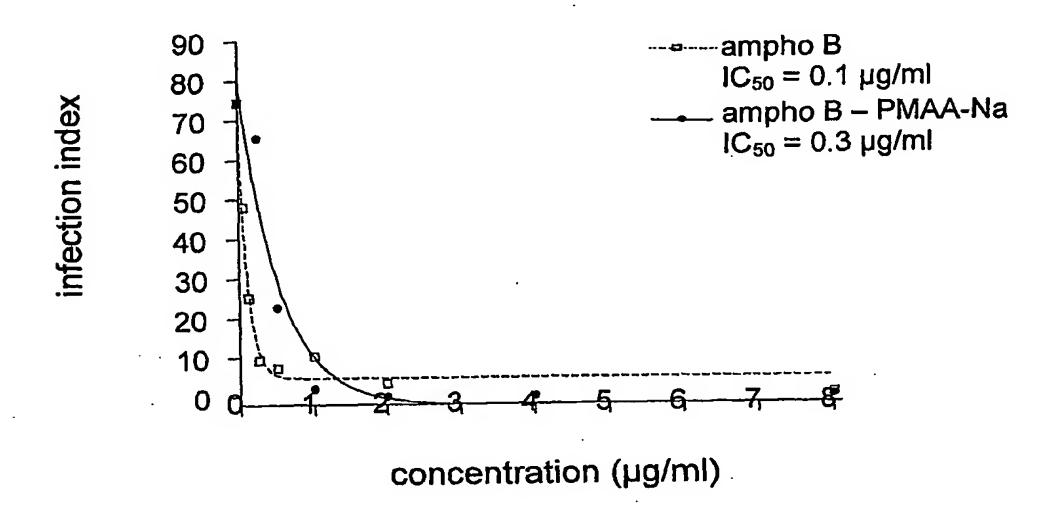


Figure 31a

Inhibition of *C. neoformans* var *neoformans* NCPF 3003 growth in monocyte derived macrophages after 3 days

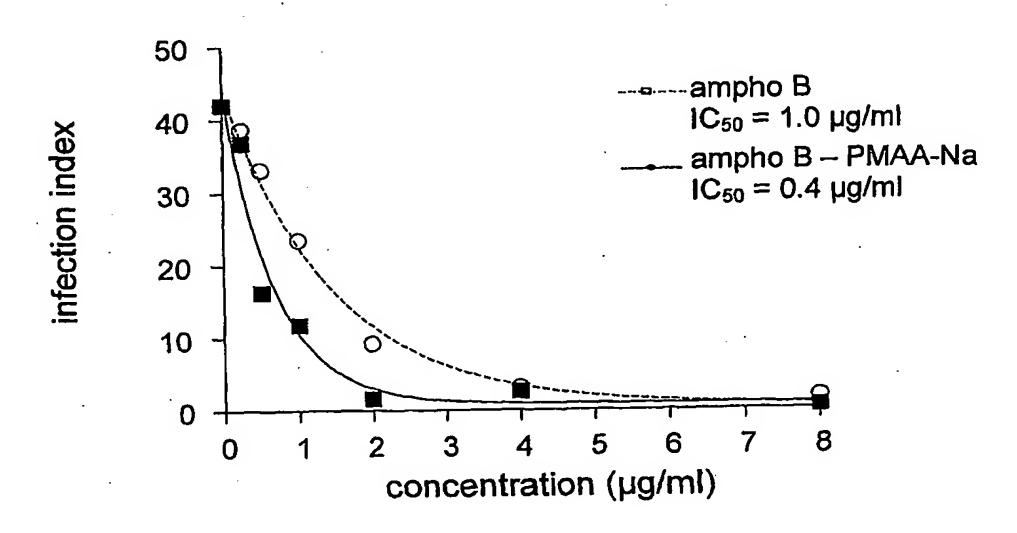


Figure 31b

Figure 31 cont.

Inhibition of *C. neoformans* var *gattii* clinical isolate growth in peritoneal macrophages after 3 days

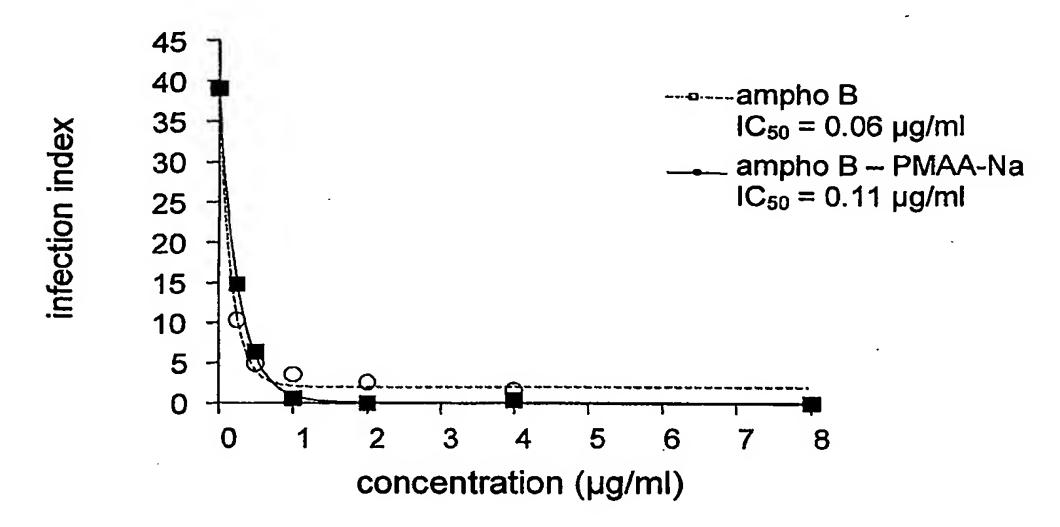


Figure 31c

Inhibition of *C. neoformans* var *gattii* clinical in monocytes derived macrophages after 3 days

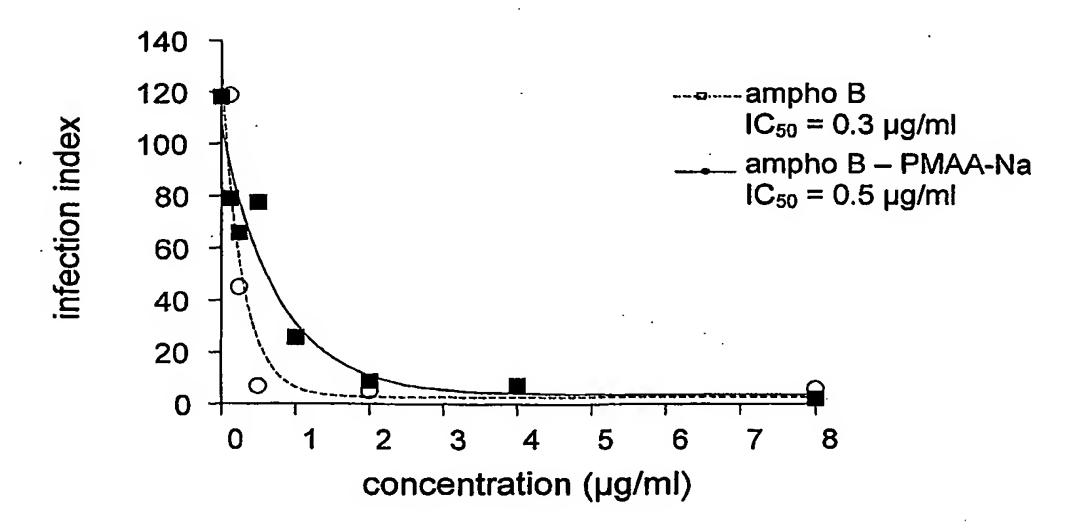


Figure 31d

Figure 32



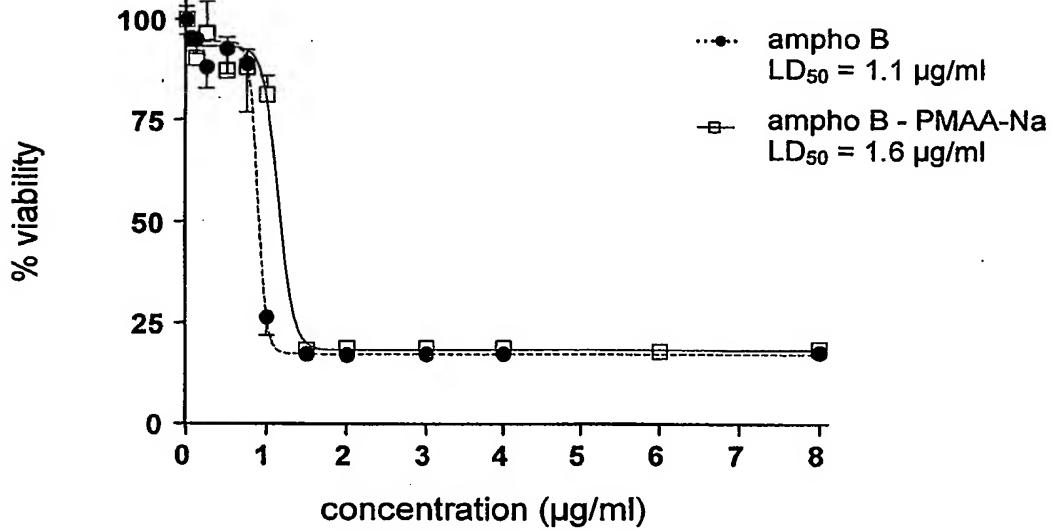


Figure 32a

Viability of *C.neoformans* var *neoformans* clinical isolate after 3 days

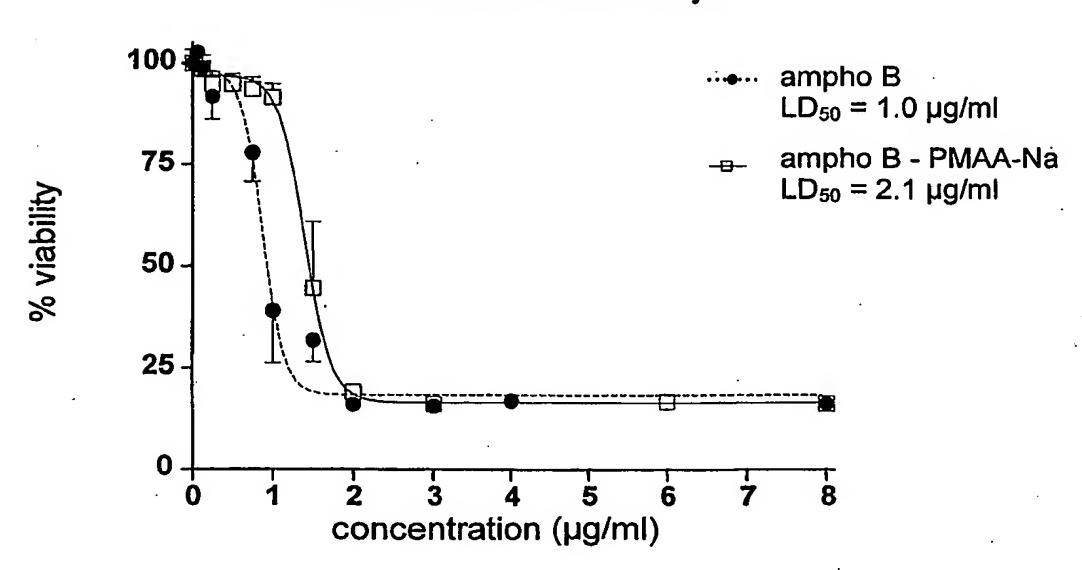


Figure 32b

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Figure 32 cont.

Viability of *C.neoformans* var *gattii* NCPF 3216 after 3 days

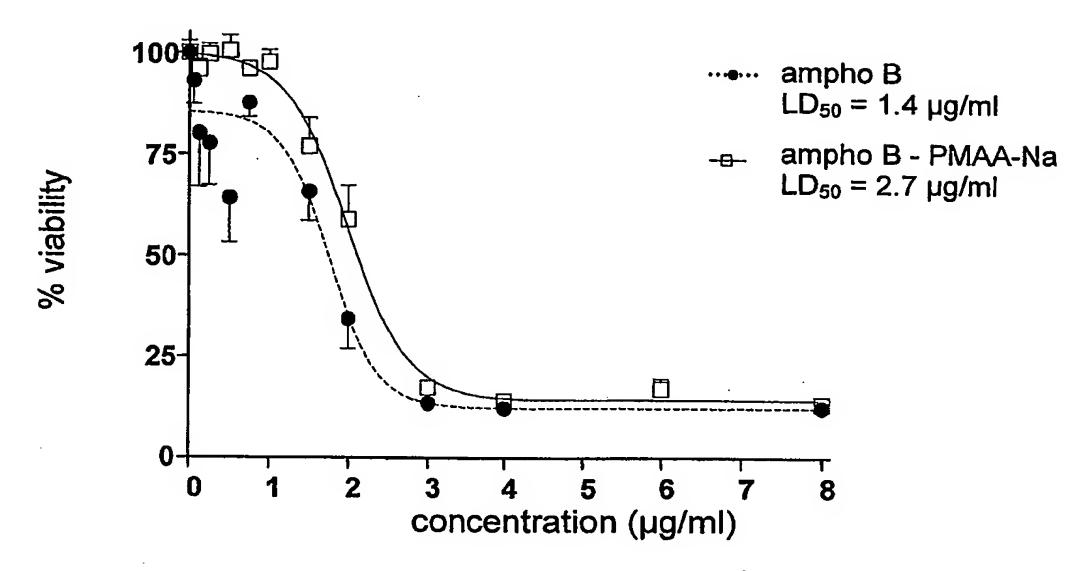


Figure 32c

Viability of *C.neoformans* var *gattii* clinical isolate after 3 days

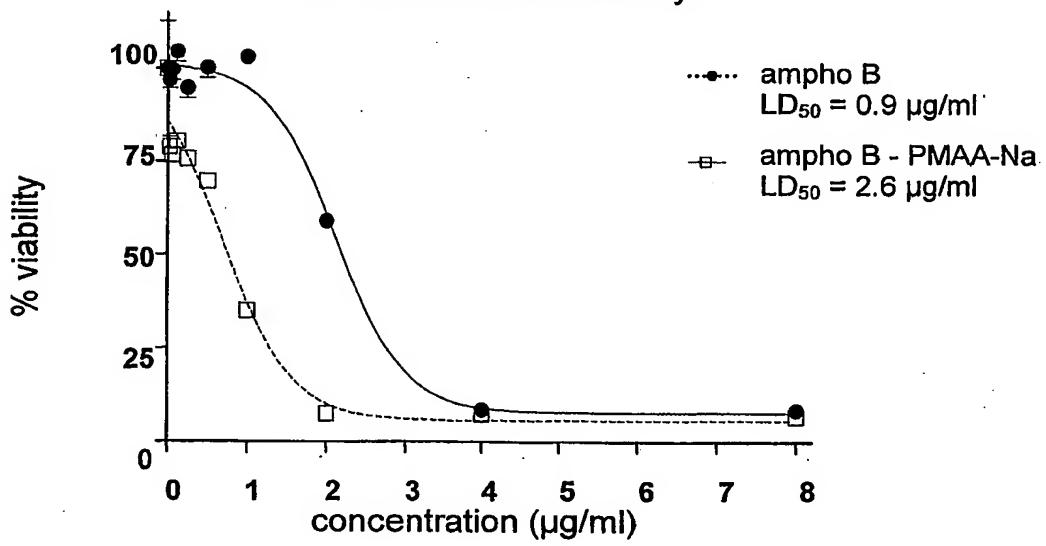


Figure 32d

Figure 33

Viability of C.albicans ATCC 90028 after 1 day

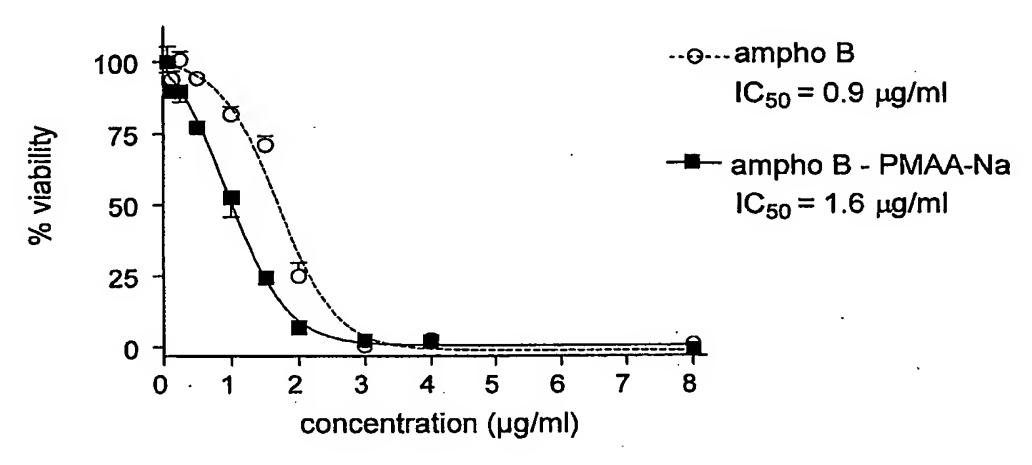


Figure 33a

Viability of C. glabrata ATCC 90030 after 1 day incubation

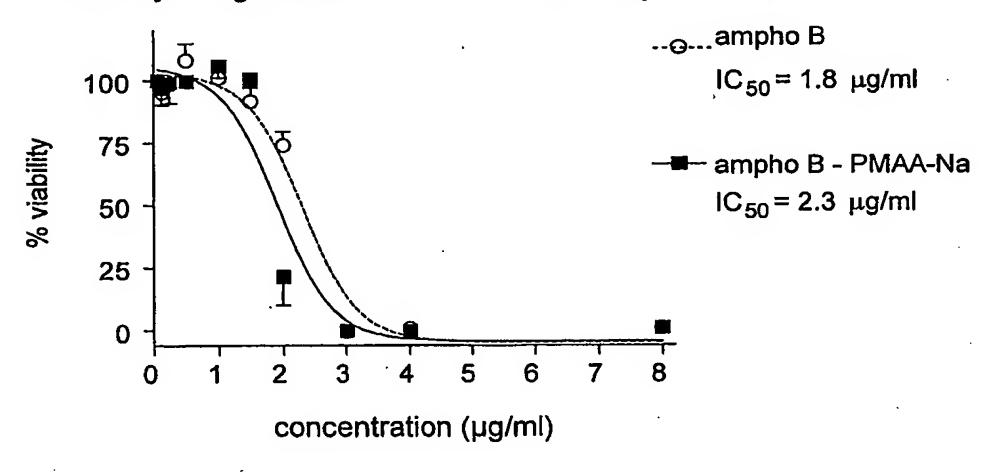


Figure 33b